



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification⁶ : A61K 48/00, 49/00, A01N 63/00	A1	(11) International Publication Number: WO 96/22793 (43) International Publication Date: 1 August 1996 (01.08.96)
(21) International Application Number: PCT/US96/01448 (22) International Filing Date: 25 January 1996 (25.01.96) (30) Priority Data: 381,603 27 January 1995 (27.01.95) US (71) Applicant: UNIVERSITY OF PITTSBURGH OF THE COMMONWEALTH SYSTEM OF HIGHER EDUCATION [US/US]; 911 William Pitt Union, Pittsburgh, PA 15260 (US). (72) Inventors: GLORIOSO, Joseph, C.; 202 Scarborough Drive, Cheswick, PA 15024 (US). EVANS, Christopher, H.; 111 Maple Terrace, Pittsburgh, PA 15211 (US). ROBBINS, Paul, D.; 527 Overlook, Pittsburgh, PA 15216 (US). (74) Agent: GOULD, Lewis, F., Jr.; Eckert Seamans Cherin & Mellott, Suite 3232, 1700 Market Street, Philadelphia, PA 19103 (US).		(81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: GENE TRANSFER FOR TREATING A CONNECTIVE TISSUE OF A MAMMALIAN HOST (57) Abstract <p>The subject invention concerns a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host including employing recombinant techniques to produce a DNA vector molecule which contains the gene encoding for the product and infecting the connective cell of the mammalian host using the DNA vector molecule using the gene coding for the product. A method is provided for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host employing non-viral means. A method to produce an animal model for the study of connective tissue pathology is also disclosed. Additionally, this invention provides a method of using <i>in vivo</i> a gene encoding an extracellular interleukin-1 binding domain of an interleukin-1 receptor.</p>		

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5 **GENE TRANSFER FOR TREATING A CONNECTIVE TISSUE
 OF A MAMMALIAN HOST**

CROSS-REFERENCE TO RELATED APPLICATION

 This is a continuation-in-part application of U.S. Application Serial No. 08/027,750,
10 filed March 8, 1993, now pending, and a continuation-in-part application of U.S. Application
Serial No. 08/163,563, filed January 18, 1994, which is a continuation application of U.S.
Application No. 07/963,928, filed October 20, 1992, now abandoned, which was a
continuation application of U.S. Application Serial No. 07/630,981, filed
December 20, 1990, now abandoned.

15 **BACKGROUND OF THE INVENTION**

Field Of The Invention

 The present invention relates to a method of introducing at least one gene encoding a
product into at least one cell of a connective tissue of a mammalian host for use in treating
the mammalian host. This method discloses employing DNA vector molecules containing a
20 gene encoding the product and infecting the connective tissue cells of the mammalian host
using the DNA vector molecule. This invention provides a method of introducing at least
one gene encoding a product into at least one cell of a connective tissue of a mammalian host
for use in treating the mammalian host including employing non-viral means for effecting
such introduction.

25 The present invention discloses *ex vivo* and *in vivo* techniques for delivery of a DNA
sequence of interest to the connective tissue cells of the mammalian host. The *ex vivo*
technique involves prior removal and culture of target autologous connective tissue cells,
in vitro infection of the DNA sequence, DNA vector or other delivery vehicle of interest into
the connective tissue cells, followed by transplantation to the modified connective tissue cells
30 to the target joint of the mammalian host, so as to effect *in vivo* expression of the gene
product of interest. The *in vivo* technique bypasses the requirement for *in vitro* culture of
target connective tissues cells; instead relying on direct transplantation of the DNA sequence,

DNA vector or other delivery vehicle to the target *in vivo* connective tissue cells, thus effecting expression of the gene product of interest.

The present invention also relates to a method to produce an animal model for the study of connective tissue pathologies and systemic indices of inflammation.

5 The present invention further relates to a method of using a gene encoding a truncated interleukin-1 receptor to resist the deleterious pathological changes associated with arthritis. More specifically, this invention provides a method wherein a gene coding for an extracellular interleukin-1 binding domain of an interleukin-1 receptor is introduced into synovial cells of a mammalian host *in vivo* for neutralizing the destructive activity of
10 interleukin-1 upon cartilage and other soft tissues. As an alternative, the patients own synovial cells are transduced *in vitro* and introduced back into the affected joint, using transplantation procedures such as for example, intra-articular injection.

As an alternative to the *in vitro* manipulation of synovia, the gene encoding the product of interest is introduced into liposomes and injected directly into the area of the
15 joint, where the liposomes fuse with synovial cells, resulting in an *in vivo* gene transfer to synovial tissue. As an additional alternative to the *in vitro* manipulation of synovia, the gene encoding the product of interest is introduced into the area of the joint as naked DNA. The naked DNA enters the synovial cell, resulting in an *in vivo* gene transfer to synovial tissue.

As an another alternative, hematopoietic progenitor cells or the mature lymphoid or
20 myeloid cells may be transfected *in vitro*, recovered and injected into the bone marrow of the patient using techniques known to the skilled artisan.

Brief Description Of The Related Art

Arthritis involves inflammation of a joint that is usually accompanied by pain and frequently changes in structure. Arthritis may result from or be associated with a number of
25 conditions including infection, immunological disturbances, trauma and degenerative joint diseases such as, for example, osteoarthritis. The biochemistry of cartilage degradation in joints and cellular changes have received considerable investigation.

In a healthy joint, cells in cartilage (chondrocytes) and the surrounding synovium (synoviocytes) are in a resting state. In this resting state, these cells secrete basal levels of

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prostaglandin E₂ and various neutral proteinases, such as, for example, collagenase, gelatinase and stromelysin, with the ability to degrade cartilage. During the development of an arthritic condition, these cells become activated. In the activated state, synoviocytes and chondrocytes synthesize and secrete large amounts of prostaglandin E₂ and neutral proteinases.

In efforts to identify pathophysiologically relevant cell activators, it has been known that the cytokine interleukin-1 activates chondrocytes and synoviocytes and induces cartilage breakdown *in vitro* and *in vivo*. Additionally, interleukin-1 is a growth factor for synoviocytes and promotes their synthesis of matrix, two properties suggesting the involvement of interleukin-1 in the synovial hypertrophy that accompanies arthritis. In contrast, interleukin-1 inhibits cartilaginous matrix synthesis by chondrocytes, thereby suppressing repair of cartilage. Interleukin-1 also induces bone resorption and thus may account for the loss of bone density seen in rheumatoid arthritis. Interleukin-1 is inflammatory, serves as a growth factor for lymphocytes, is a chemotactic factor and a possible activator of polymorphonuclear leukocytes (PMNs). When present in a sufficient concentration, interleukin-1 may cause fever, muscle wasting and sleepiness.

The major source of interleukin-1 in the joint is the synovium. Interleukin-1 is secreted by the resident synoviocytes, which are joined under inflammatory conditions by macrophages and other white blood cells.

Much attention has been devoted to the development of a class of agents identified as the "Non-Steroidal Anti-Inflammatory Drugs" (hereinafter "NSAIDs"). The NSAIDs inhibit cartilage synthesis and repair and control inflammation. The mechanism of action of the NSAIDs appears to be associated principally with the inhibition of prostaglandin synthesis in body tissues. Most of this development has involved the synthesis of better inhibitors of cyclo-oxygenase, a key enzyme that catalyzes the formation of prostaglandin precursors (endoperoxides) from arachidonic acid. The anti-inflammatory effect of the NSAIDs is thought to be due in part to inhibition of prostaglandin synthesis and release during inflammation. Prostaglandins are also believed to play a role in modulating the rate and extent of leukocyte infiltration during inflammation. The NSAIDs include, such as, for example, acetylsalicylic acid (aspirin), fenoprofen calcium (Nalfon Pulvules, Dista

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Products Company), ibuprofen (Motrin , The Upjohn Company), and indomethacin (Indocin , Merck, Sharp & Dohme).

In contrast, the studies upon which the present invention is based show that production of the various neutral proteinases with the ability to degrade cartilage occurs even if prostaglandin synthesis is completely blocked.

Therapeutic intervention in arthritis is hindered by the inability to target drugs, such as the NSAIDs, to specific areas within a mammalian host, such as, for example a joint. Traditional routes of drug delivery, such as for example, oral, intravenous or intramuscular administration, depend upon vascular perfusion of the synovium to carry the drug to the joint. This is inefficient because transsynovial transfer of small molecules from the synovial capillaries to the joint space occurs generally by passive diffusion. This diffusion is less efficient with increased size of the target molecule. Thus, the access of large drug molecules, for example, proteins, to the joint space is substantially restricted. Intra-articular injection of drugs circumvents those limitations; however, the half-life of drugs administered intra-articularly is generally short. Another disadvantage of intra-articular injection of drugs is that frequent repeated injections are necessary to obtain acceptable drug levels at the joint spaces for treating a chronic condition such as, for example, arthritis. Because therapeutic agents heretofore could not be selectively targeted to joints, it was necessary to expose the mammalian host to systemically high concentrations of drugs in order to achieve a sustained, intra-articular therapeutic dose. Exposure of non-target organs in this manner exacerbated the tendency of anti-arthritis drugs to produce serious side effects, such as for example, gastrointestinal upset and changes in the hematological, cardiovascular, hepatic and renal systems of the mammalian host.

It has been shown that genetic material can be introduced into mammalian cells by chemical or biologic means. Moreover, the introduced genetic material can be expressed so that high levels of a specific protein can be synthesized by the host cell. Cells retaining the introduced genetic material may include an antibiotic resistance gene thus providing a selectable marker for preferential growth of the transduced cell in the presence of the corresponding antibiotic. Chemical compounds for inhibiting the production of interleukin-1 are also known.

U.S. Patent No. 4,778,806 discloses a method of inhibiting the production of interleukin-1 by monocytes and/or macrophages in a human by administering through the parenteral route a 2-2'-[1,3-propan-2-onediyl-bis (thio)] bis-1 H-imidazole or a pharmaceutically acceptable salt thereof. This patent discloses a chemical compound for inhibiting the production of interleukin-1. By contrast, in one embodiment of the present invention, gene therapy is employed that is capable of binding to and neutralizing interleukin-1.

U.S. Patent No. 4,780,470 discloses a method of inhibiting the production of interleukin-1 by monocytes in a human by administering a 4,5-diaryl-2 (substituted) imidazole. This patent also discloses a chemical compound for inhibiting the production of interleukin-1.

U.S. Patent No. 4,794,114 discloses a method of inhibiting the 5-lipoxygenase pathway in a human by administering a diaryl-substituted imidazole fused to a thiazole, pyrrolidine or piperidine ring or a pharmaceutically acceptable salt thereof. This patent also discloses a chemical compound for inhibiting the production of interleukin-1.

U.S. Patent No. 4,870,101 discloses a method for inhibiting the release of interleukin-1 and for alleviating interleukin-1 mediated conditions by administering an effective amount of a pharmaceutically acceptable anti-oxidant compound such as disulfiram, tetrakis [3-(2,6-di-tert-butyl-4-hydroxyphenyl) propionyloxy methyl] methane or 2,4-di-isobutyl-6-(N,N-dimethylamino methyl)-phenol. This patent discloses a chemical compound for inhibiting the release of interleukin-1.

U.S. Patent No. 4,816,436 discloses a process for the use of interleukin-1 as an anti-arthritic agent. This patent states that interleukin-1, in association with a pharmaceutical carrier, may be administered by intra-articular injection for the treatment of arthritis or inflammation. In contrast, the present invention discloses a method of using and preparing a gene that is capable of binding to and neutralizing interleukin-1 as a method of resisting arthritis.

U.S. Patent No. 4,935,343 discloses an immunoassay method for the detection of interleukin-1 beta that employs a monoclonal antibody that binds to interleukin-1 beta but does not bind to interleukin-1 alpha. This patent discloses that the monoclonal antibody

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binds to interleukin-1 beta and blocks the binding of interleukin-1 beta to interleukin-1 receptors, and thus blocking the biological activity of interleukin-1 beta. The monoclonal antibody disclosed in this patent may be obtained by production of an immunogen through genetic engineering using recombinant DNA technology. The immunogen is injected into a mouse and thereafter spleen cells of the mouse are immortalized by fusing the spleen cells with myeloma cells. The resulting cells include the hybrid continuous cell lines (hybridomas) that may be later screened for monoclonal antibodies. This patent states that the monoclonal antibodies of the invention may be used therapeutically, such as for example, in the immunization of a patient, or the monoclonal antibodies may be bound to a toxin to form an immunotoxin or to a radioactive material or drug to form a radio pharmaceutical or pharmaceutical.

U.S. Patent No. 4,766,069 discloses a recombinant DNA cloning vehicle having a DNA sequence comprising the human interleukin-1 gene DNA sequence. This patent provides a process for preparing human interleukin-1 beta, and recovering the human interleukin-1 beta. This patent discloses use of interleukin-1 as an immunological reagent in humans because of its ability to stimulate T-cells and B-cells and increase immunoglobulin synthesis.

U.S. Patent No. 4,396,601 discloses a method for providing mammalian hosts with additional genetic capability. This patent provides that host cells capable of regeneration are removed from the host and treated with genetic material including at least one marker which allows for selective advantage for the host cells in which the genetic material is capable of expression and replication. This patent states that the modified host cells are then returned to the host under regenerative conditions. In the present invention, genetic material may be directly introduced (a) into host cells *in vivo* or (b) into synoviocytes *in vitro* for subsequent transplantation back into the patient's joints.

U.S. Patent No. 4,968,607 discloses a DNA sequence encoding a mammalian interleukin-1 receptor protein which exhibits interleukin-1 binding activity.

U.S. Patent No. 5,081,228 discloses a DNA sequence encoding both the murine and human interleukin-1 receptor. This patent also provides a process for the *in vitro* expression of said DNA sequences.

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U.S. Patent No. 5,180,812 discloses a substantially pure preparation of the human interleukin-1 receptor protein.

In spite of these prior art disclosures, there remains a very real and substantial need for a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host *in vitro*, or alternatively *in vivo*, for use in treating the mammalian host. Further, there is a need for a process wherein a gene encoding a truncated interleukin-1 receptor is used to resist the deleterious pathological changes associated with arthritis. More specifically there is a need for such a process where a gene coding for the extracellular interleukin-1 binding domain of the interleukin-1 receptor, capable of binding to and neutralizing interleukin-1 is expressed in host synovial cells *in vivo*. There is also a need to utilize one or more additional DNA sequences for delivery to and expression of a protein or protein fragment within a target host connective tissue cell, such as a synovial cell, so as to effect a treatment of various joint pathologies and concomitant systemic indices of inflammation.

SUMMARY OF THE INVENTION

The present invention has met the hereinbefore described need. A method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host is provided for in the present invention. This method includes employing recombinant techniques to produce a DNA vector molecule containing the gene encoding for the product and infecting the connective tissue cell of the mammalian host using the DNA vector molecule containing the gene coding for the product. The DNA vector molecule can be any DNA molecule capable of being delivered and maintained within the target cell or tissue such that the gene encoding the product of interest can be stably expressed. The DNA vector molecule preferably utilized in the present invention is either a viral DNA vector molecule or a plasmid DNA viral molecule. This method preferably includes introducing the gene encoding the product into the cell of the mammalian connective tissue for a therapeutic use.

One *ex vivo* method of treating a connective tissue disorder disclosed throughout this specification comprises initially generating a recombinant viral vector which contains a DNA sequence encoding a protein or biologically active fragment thereof. This recombinant viral

vector is then used to infect a population of *in vitro* cultured connective tissue cells, resulting in a population of transfected connective cells. These transfected connective tissue cell are then transplanted to a target joint space of a mammalian host, effecting subsequent expression of the protein or protein fragment within the joint space. Expression of this DNA sequence of interest is useful in substantially reducing at least one deleterious joint pathology or indicia of inflammation normally associated with a connective tissue disorder.

The connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament, preferably synovial cells.

It is also preferred that a retroviral vector, such as MFG, be utilized as the viral vector.

Another preferred step in this *ex vivo* method is transplantation of transduced cells by intraarticular injection.

It will be understood by the artisan of ordinary skill that the preferred source of cells for treating a human patient are the patients own cells, such as autologous synovial cells.

More specifically, this method includes employing as the gene a gene capable of encoding at least one of the materials which is selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or a biologically active derivative or fragment thereof, preferably MFG-IRAP, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or a biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or a biologically active derivative or fragment thereof, (d) a soluble TNF- α receptor protein or a biologically active derivative or fragment thereof; (e) a proteinase inhibitor, and (f) a cytokine, and employing as the viral vector at least one vector which is selected from the group which includes (a) a retroviral vector including at least one of the materials selected from the group which includes MFG and BAG, (b) an adeno-associated virus, (c) an adenovirus, and (d) a herpes virus, including but not limited to herpes simplex 1 or herpes simplex 2.

A further embodiment of the present invention includes employing as the gene a gene capable of encoding at least one of the materials which is selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or a biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a

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beta-galactosidase protein or a biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or a biologically active derivative or fragment thereof, (d) a soluble TNF- α receptor protein or a biologically active derivative or fragment thereof; (e) a proteinase inhibitor, and (f) a cytokine, and employing as the DNA plasmid vector any DNA plasmid vector known to one of ordinary skill in the art capable of stable maintenance within the targeted cell or tissue upon delivery, regardless of the method of delivery utilized. One such method is the direct delivery of the DNA vector molecule, whether it be a viral or plasmid DNA vector molecule, to the target cell or tissue. This method also includes employing as the gene a gene capable of encoding at least one of the materials selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or biologically active derivative or fragment thereof, (d) a soluble TNF- α receptor protein or a biologically active derivative or fragment thereof; (e) a proteinase inhibitor, and (f) a cytokine. In a specific method disclosed as an example, and not as a limitation to the present invention, a DNA plasmid vector containing the interleukin-1 beta (IL-1 β) coding sequence was ligated downstream of the cytomegalovirus (CMV) promoter. This DNA plasmid construction was encapsulated within liposomes and injected intra-articularly into the knee joints of recipient rabbits. IL-1 β was expressed and significant amounts of interleukin-1 beta was recovered from the synovial tissue. An alternative is injection of the naked plasmid DNA into the knee joint, allowing direct transfection of the DNA into the synovial tissue. Injection of IL-1 β into the joint of a mammalian host allows for prolonged study of various joint pathologies and systemic indices of inflammation, as described within this specification.

Another embodiment of this invention provides a method for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host. This method includes employing non-viral means for introducing the gene encoding for the product into the connective tissue cell. More specifically, this method includes employing non-viral means which is selected from at least one of the group which includes (a) at least one liposome, (b) $\text{Ca}_3(\text{PO}_4)_2$, (c) electroporation,

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and (d) DEAE-dextran, and includes employing as the gene a gene capable of encoding at least one of the materials selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or biologically active derivative or fragment thereof, (d) a soluble TNF- α receptor protein or a biologically active derivative or fragment thereof; (e) a proteinase inhibitor, and (f) a cytokine.

A further embodiment of this invention provides an additional method for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host. This additional method includes employing the biologic means of utilizing a virus to deliver the DNA vector molecule to the target cell or tissue. Preferably, the virus is a psuedovirus, the genome having been altered such that the psuedovirus is capable only of delivery and stable maintenance within the target cell; but not retaining an ability to replicate within the target cell or tissue. The altered viral genome is further manipulated by recombinant DNA techniques such that the viral genome acts as a DNA vector molecule which contains the heterologous gene of interest to be expressed within the target cell or tissue. This method also includes employing as the gene a gene capable of encoding at least one of the materials selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or biologically active derivative or fragment thereof, (d) a soluble TNF- α receptor protein or a biologically active derivative or fragment thereof, (e) a proteinase inhibitor, and (f) a cytokine.

A further embodiment of this invention provides for an animal model to study connective tissue pathologies and indices of systemic inflammation. This model utilizes either *ex vivo* or *in vivo* delivery of at least one gene or DNA sequence of interest encoding a product into at least one cell of a connective tissue of a mammalian host. Examples of joint pathologies which can be studied in the present invention include, but are by no means limited to, joint pathologies such as leukocytosis, synovitis, cartilage breakdown and

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suppression of cartilage matrix synthesis. Examples of indices of systemic inflammation which include, but are by no means limited to, erythrocyte sedimentation rate, fever and weight loss.

5 An embodiment of the present invention is a method to produce an animal model for the study of joint pathologies. This embodiment comprises generating a recombinant viral vector which contains a DNA sequence encoding a protein or biologically active fragment thereof, infecting a population of *in vitro* cultured connective tissue cells with said recombinant viral vector, resulting in a population of transfected connective cells; and, transplanting said transfected connective cells to a joint space of a mammalian host, such that
10 subsequent expression of said protein within said joint space substantially reduces at least one deleterious joint pathology or indicia of inflammation normally associated with a connective tissue disorder.

The connective tissue cells from this embodiment are also selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament, preferably
15 synovial cells, including but not limited to autologous cells removed directly from the mammalian host of which the target joint resides.

As with other *ex vivo* based methods disclosed throughout this specification, a preferable method of transplantation of transduced connective cells, such as synovial cells, is by intraarticular injection.

20 A DNA sequence exemplified for animal model studies is a DNA sequence encoding encoding human IL-1 α or a biologically active fragment thereof.

Another DNA sequence exemplified for animal model studies is a DNA sequence encoding human tumor necrosis factor- α or a biologically active fragment thereof.

Another embodiment of a method to produce an animal model for the study of joint
25 pathologies utilizes a recombinant DNA plasmid vector, which contains the DNA sequence of interest encoding a protein or biologically active fragment thereof. This recombinant DNA plasmid vector is used to transform a population of *in vitro* cultured connective tissue cells. The transformed connective cells, preferably synovial cells, are transplanted to a joint space of a mammalian host, so as to provide data regarding various joint pathologies and systemic
30 indices of inflammation associated with connective tissue disorders.

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This particular embodiment is exemplified by the *ex vivo* based delivery of MFG-IL-1 β to a target rabbit knee joint, causing various joint pathologies and systemic indices of inflammations.

Another exemplification of this particular embodiment of the present invention is delivery of the CMV-IL-1 β plasmid construction to the rabbit knee joint via liposome-mediated delivery.

An animal model as described and exemplified in this specification measures the ability of various gene therapy applications disclosed throughout this specification to withstand challenges from known causative agents (such as IL-1 β) of joint pathologies and inflammatory side effects.

An additional embodiment of the present invention relates to a method of using a DNA sequence encoding a biologically active interleukin-1 receptor antagonist (IRAP) or portion thereof for treatment of connective tissue joint pathologies. The DNA sequence encoding IRAP or a biologically active fragment thereof may be delivered to the connective tissue of a mammalian host by any combination of various vector strategies and transduction techniques disclosed throughout this specification.

A preferred method of the embodiment of delivering IRAP to a target joint space involves delivery of the IPAP gene to the synovial lining of a mammalian host through use of a retroviral vector with the *ex vivo* technique disclosed within this specification. In other words, a DNA sequence of interest encoding a functional IPAP protein or protein fragment is subcloned into a retroviral vector of choice, the recombinant viral vector is then grown to adequate titers and used to infect *in vitro* cultured synovial cells, and the transduced synovial cells, preferably autografted cells, are transplanted into the joint of interest, preferably by intra-articular injection.

Another preferred method of the present invention involves direct *in vivo* delivery of the IRAP gene to the synovial lining of a mammalian host through use of either an adenovirus vector, adeno-associated virus (AAV) vector or herpes-simples virus (HSV) vector. In other words, a DNA sequence of interest encoding a functional IRAP protein or protein fragment is subcloned into the respective viral vector, the IRAP containing viral vector is then grown to adequate titers and directed into the joint space, preferably by

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intra-articular injection. A retroviral-IRAP construct, such as MFG-IRAP may also be utilized to directly target previously inflamed connective tissue cells within the joint space.

Direct intra-articular injection of a DNA molecule containing the gene of interest into the joint results in transfection of the recipient synovial cells and hence bypasses the requirement of removal, *in vitro* culturing, transfection, selection, as well as transplanting the DNA vector containing - synoviocytes (as disclosed in the Example section) to promote stable expression of the heterologous gene of interest. Methods of presenting the DNA molecule to the target connective tissue of the joint includes, but is not limited to, encapsulation of the DNA molecule into cationic liposomes, subcloning the DNA sequence of interest in a retroviral vector as described throughout this specification, or the direct injection of the DNA molecule itself into the joint. The DNA molecule, regardless of the form of presentation to the knee joint, is preferably presented as a DNA vector molecule, either as recombinant viral DNA vector molecule or a recombinant DNA plasmid vector molecule. Expression of the heterologous gene of interest is ensured by inserting a promoter fragment active in eukaryotic cells directly upstream of the coding region of the heterologous gene. One of ordinary skill in the art may utilize known strategies and techniques of vector construction to ensure appropriate levels of expression subsequent to entry of the DNA molecule into the synovial tissue. *In vivo* delivery of various viral and non-viral vectors to the rabbit knee joint are described in Example XIV.

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of the human TNF- α soluble receptor or a biologically active fragment thereof.

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various cytokines which possess anti-inflammatory and immunomodulatory characteristics, including but by no means limited to interleukin-4 and interleukin-10.

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within

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this specification so as to effect *in vivo* expression of various antiadhesion molecules so as to inhibit cell-cell and cell-matrix interactions. Examples of such proteins or protein fragments include but are not limited to soluble ICAM-1 and soluble CD44.

5 Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various cartilage growth factors, including but not limited to IGF-1 and TGF- β .

10 Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various free radical antagonists, thus preventing the deleterious effects of free radical formation within the afflicted joint. Examples include but are not limited to the superoxide dismutase and proteins or protein fragments which inhibit NO.

15 Another embodiment of the present invention regarding delivery of the IRAP gene to the synovial lining of a mammalian host involves use subcloning this DNA sequence of interest into a viral vector such as adenovirus, adeno-associated virus and herpes-simplex virus. The respective recombinant IRAP based viral vector is then delivered to the joint by direct *in vivo* injection so as to effect *in vivo* expression of the IRAP protein or biologically active fragment thereof.

20 Another embodiment of this invention provides a method of using the gene encoding an extracellular interleukin-1 binding domain of the interleukin-1 receptor. This gene is capable of binding to and neutralizing interleukin-1 *in vivo* to substantially resist the degradation of cartilage in a mammalian host. Unlike previous pharmacological efforts, the method of this invention employs gene therapy *in vivo* to address the chronic debilitating effects of
25 arthritis.

A preferred method of using the gene coding for the truncated interleukin-1 receptor of this invention involves employing recombinant techniques to generate a cell line which produces infectious retroviral particles containing the gene coding for the truncated interleukin-1 receptor. The producer cell line is generated by inserting the gene coding into

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a retroviral vector under the regulation of a suitable eukaryotic promoter, transfecting the retroviral vector containing the gene coding into the retroviral packaging cell line for the production of a viral particle that is capable of expressing the gene coding for the truncated interleukin-1 receptor, and infecting the synovial cells of a mammalian host using the viral particle.

More specifically, the method of using the hereinbefore described gene coding for the truncated interleukin-1 receptor involves introducing the viral particles obtained from the retroviral packaging cell line directly by intra-articular injection into a joint space of a mammalian host that is lined with synovial cells. In a preferred embodiment, synoviocytes recovered from the knee joint are cultured *in vitro* for subsequent utilization as a delivery system for gene therapy. It will be apparent that Applicants are not limited to the use of the specific synovial tissue disclosed. It would be possible to utilize other tissue sources, such as skin cells, for *in vitro* culture techniques. The method of using the gene of this invention may be employed both prophylactically and in the therapeutic treatment of arthritis. It will also be apparent that Applicants are not limited to prophylactic or therapeutic applications in treating only the knee joint. It would be possible to utilize the present invention either prophylactically or therapeutically to treat arthritis in any susceptible joint.

In another embodiment of this invention, a method of using the hereinbefore described gene coding for the truncated interleukin-1 receptor involves infecting synovial cells in culture with the viral particles and subsequently transplanting the infected synovial cells back into the joint. This method of using the gene of this invention may also be employed prophylactically and in the therapeutic treatment of arthritis in any area susceptible to the disorder.

In another embodiment of this invention, a method of using the gene coding for an extracellular interleukin-1 binding domain of the interleukin-1 receptor that is capable of binding to and neutralizing interleukin-1 includes employing recombinant techniques to produce a retrovirus vector carrying two genes. The first gene encodes the extracellular interleukin-1 binding domain of the interleukin receptor, and the second gene encodes for selectable antibiotic resistance. This method of use involves transfecting the retrovirus vector

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into a retrovirus packaging cell line to obtain a cell line producing infectious retroviral particles carrying the gene.

Another embodiment of this invention provides a method of preparing a gene encoding an extracellular interleukin-1 binding domain of the interleukin-1 receptor including synthesizing the gene by a polymerase chain reaction, introducing the amplified interleukin-1 receptor coding sequence into a retroviral vector, transfecting the retroviral vector into a retrovirus packaging cell line and collecting viral particles from the retrovirus packaging cell line.

In another embodiment of this invention, a compound for parenteral administration to a patient in a therapeutically effective amount is provided for that contains a gene encoding an extracellular interleukin-1 binding domain of the interleukin-1 receptor and a suitable pharmaceutical carrier.

Another embodiment of this invention provides for a compound for parenteral administration to a patient in a prophylactically effective amount that includes a gene encoding an extracellular interleukin-1 binding domain of the interleukin-1 receptor and a suitable pharmaceutical carrier.

An additional embodiment of the invention involves transfection of hematopoietic progenitor cells or mature lymphoid or myeloid cells with a DNA vector molecule containing any of the gene or genes disclosed throughout the specification. The transfected cells are recovered and injected into the bone marrow of the patient using techniques known and available to one of ordinary skill in the art. It will be possible, within the scope of this method, to use cells derived from donor bone marrow instead of cells derived from recipient bone marrow so as to modify rejection.

It is an object of the present invention to provide a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host.

It is an object of the invention to provide a method of introducing a gene encoding a product into at least one cell of a connective tissue of a mammalian host for a therapeutic use.

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It is an object of the present invention to provide a method of introducing into the synovial lining cells of a mammalian arthritic joint at least one gene which codes for proteins having therapeutic properties.

5 It is an object of the present invention to provide an animal model for the study of connective tissue pathology.

It is an object of the present invention to provide a method of using *in vivo* a gene coding for the extracellular interleukin-1 binding domain of the interleukin-1 receptor that is capable of binding to and neutralizing substantially all isoforms of interleukin-1, including interleukin-1 alpha and interleukin-1 beta.

10 It is an object of the present invention to provide a method of using *in vivo* a gene coding for IRAP or a biologically active derivative thereof which is a competitive inhibitor of and therefore substantially neutralizes all isoforms of interleukin-1, including interleukin-1 alpha and interleukin-1 beta.

15 It is an object of the present invention to provide a method of using a gene *in vivo* in a mammalian host that is capable of binding to and neutralizing substantially all isoforms of interleukin-1 and thus, substantially resist the degradation of cartilage and protect surrounding soft tissues of the joint space.

20 It is an object of the present invention to provide a method of using *in vivo* a gene coding for the extracellular interleukin-1 binding domain of the interleukin-1 receptor that is capable of binding to and neutralizing substantially all isoforms of interleukin-1 for the prevention of arthritis in patients that demonstrate a high susceptibility for developing the disease.

25 It is an object of the present invention to provide a method of using *in vivo* a gene coding for IRAP that is capable of acting as a competitive inhibitor of and therefore substantially neutralizes all isoforms of interleukin-1 for the prevention of arthritis in patients that demonstrate a high susceptibility for developing the disease.

30 It is an object of the present invention to provide a method of using *in vivo* a gene coding for an extracellular interleukin-1 binding domain of an interleukin-1 receptor that is capable of binding to and neutralizing substantially all isoforms of interleukin-1 for the treatment of patients with arthritis.

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It is an object of the present invention to provide a method of using *in vivo* a gene coding for IRAP or a biologically active derivative thereof which is a competitive inhibitor of and therefore substantially neutralizes all isoforms of interleukin-1 for the treatment of patients with arthritis.

5 It is an object of the present invention to provide a method of using *in vivo* a gene or genes that address the chronic debilitating pathophysiology of arthritis.

It is a further object of the present invention to provide a compound for parenteral administration to a patient which comprises a gene encoding an extracellular interleukin-1 binding domain of the interleukin-1 receptor and a suitable pharmaceutical carrier.

10 It is a further object of the present invention to provide a compound for parenteral administration to a patient which comprises a gene encoding IRAP and a suitable pharmaceutical carrier.

These and other objects of the invention will be more fully understood from the following description of the invention, the referenced drawings attached hereto and the claims
15 appended hereto.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows the structure of the cDNA encoding the human interleukin-1 receptor antagonist protein (IRAP) gene inserted into the NcoI and BamHI cloning sites of the retroviral vector MFG.

20 Figure 2 shows the structure of the cDNA encoding the human interleukin-1 receptor antagonist protein (IRAP) gene with a selectable neo marker inserted into the retroviral vector MFG.

Figure 3 shows a micrograph of synovium recovered from the knee of a rabbit approximately one month after intra-articular injection of Lac Z⁺, neo synoviocytes
25 employing the methods of this invention.

Figure 4 shows a Western blot demonstrating the production of interleukin-1 receptor antagonist protein by four cultures of HIG-82 cells (Georgescu, et al., 1988, In Vitro 24: 1015-1022) infected using the method of this invention employing the MFG-IRAP viral vector.

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Figure 5 shows data demonstrating the inhibition of chondrocytes by the addition of medium conditioned by MFG-IRAP infected HIG-82 cells.

Figure 6 shows the uptake and expression of the Lac Z gene by synoviocytes using lipofection. Well 1 - Control cells, treated with liposomes alone; Well 2 - Control cells, treated with DNA alone; Well 3 - DNA + 150 nmole liposomes; Well 4 - DNA + 240 nmole liposomes; Well 5 - DNA + 300 nmole liposomes; Well 6 - DNA + 600 nmole liposomes.

Figure 7 shows the interleukin-1 binding domain amino acid arrangement.

Figures 8A-8C show the amino acid and nucleotide sequence of the human (SEQ ID NOS 1 and 2) and mouse (SEQ ID NOS 3 and 4) interleukin-1 receptors.

Figure 9 shows gene encoding a truncated interleukin-1 receptor inserted into a retroviral vector.

Figure 10 shows antiinflammatory properties of the MFG-IRAP transgene. MFG-IRAP/HIG-82 cells (10^7) or untransduced HIG-82 cells (10^7) were transplanted to the knee joints of rabbits 3 days before intraarticular challenge with the indicated amounts of recombinant human interleukin-1 beta (rhIL-1 β). Lavage of joints occurred 18 hours later, after which infiltrating leukocytes were counted.

Figure 11 shows levels of human IRAP in rabbit knees four days following transplant of synoviocytes. Either untransduced (naive) HIG-82 cells or cells carrying a human IRAP gene (MFG-IRAP/HIG-82) were injected intra-articularly in the knee joints of rabbits (10^7 cells/knee). Four days later, knees were lavaged and the concentration of human IRAP determined by ELISA. Values given are means + S.D. (n=15).

Figure 12 shows inhibition of IL-1 induced leukocyte infiltration in knees expressing IRAP gene. Either naive or IRAP-transduced HIG-82 cells were transplanted into rabbit knee joints, as indicated. Three days later 0-100 pg/knee hrIL-1 β was intra-articularly injected at the indicated doses. The following day, knee joints were lavaged and the leukocytic infiltrate analyzed by counting with a hemocytometer and by cytopinning. Means + S.E. (n=3). (a) White blood cells (WBC) per knee. (b) Stained cytospin preparation of lavages from control knee injected with IL-1. Preparation was diluted 1:10 prior to

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cytospinning. (c) Stained cytospin preparation of lavages from IRAP-secreting knee injected with IL-1. The preparation was not diluted.

Figure 13 shows suppression of IL-1 induced loss of proteoglycans from articular cartilage. Either naive or IRAP-transduced HIG-82 cells were transplanted into rabbits knee joints. Three days later, 0-200 pg/knee hrIL-1 was intra-articularly injected at the indicated doses. The following day, knee joints were lavaged and the level of glycosaminoglycans (GAG) as an index of cartilage breakdown was determined.

Figure 14A-D shows suppression of IL-1 mediated synovial changes in knees expressing IRAP. Ten pg hrIL-1B was injected intra-articularly in each case. Synovia were harvested 18 hours after injection of IL-1B, i.e. 4 days after transplantation of naive or IRAP-secreting HIG-82 cells. (a) Control synovium following injection of IL-1, magnification x 10. (b) IRAP-secreting synovium following injection of IL-1, magnification x 10. (c) Control synovium following injection of IL-1, magnification x 160. (d) IRAP-secreting synovium, magnification x 160.

Figure 15 shows expression of human IRAP in normal and arthritic knees of rabbits. Antigen-induced arthritis was initiated by injecting 5mg ovalbumin into one knee joint (arthritic knee) of pre-sensitized rabbits on day 1. The contralateral knee (non-arthritic knee) received carrier solution only. On day 2, autologous synoviocytes (10^7 /knee in 1ml saline) were transferred to selected knee joints by intraarticular injection. Certain non-arthritic knees and arthritic knees received cells transduced with the human IRAP gene. Other non-arthritic and arthritic knees received untransduced cells or cells transduced with lac Z and neo^r genes (controls). As the results obtained with these two types of control cells were indistinguishable, they have been pooled in the figures. Detailed methods for synoviocyte culture, transduction and intraarticular implantation are disclosed throughout this specification

On day 4, knees were lavaged with 1 ml saline. On day 7, rabbits were killed and the knees again lavaged. The concentrations of human IRAP in the lavage fluids were determined by ELISA using a commercial kit (R&D Systems, Minneapolis, MN). Values given are means \pm S.E. Numbers of knees are shown above each column. Asterisks denote values which differ at $p < 0.05$ (t-test).

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Figure 16 shows concentrations of rabbit IL-1 β in the normal and arthritic knee joints of rabbits. Experimental conditions were identical to those described in Figure 15. However, lavage fluids were assayed for rabbit IL-1 α and rabbit IL-1 β by RIA using a commercial kit (Cytokine Sciences, Boston, MA). Low levels of IL-1 β are present in non-arthritic knees as a reflection of the slight inflammatory effects provoked by intraarticular injection. No IL-1 α was detectable in any of the samples. Values given are means \pm S.E. Numbers of knees are shown above each column. Asterisks denote values which differ at $p < 0.05$ (t-test).

Figure 17A-B shows the effect of IRAP gene transfer on cartilage matrix metabolism. Experimental conditions were as described for Figure 15, except that rabbits were killed both at days 4 and 7. GAG concentrations in the lavage fluids (Figure 17a) were measured spectrophotometrically by the dimethylmethylen blue assay (Farndale, et al., 1986, Biochim. Biophys. Acta 883: 173-177). Fragments of articular cartilage were shaved from the femoral condyles of the knees and GAG synthesis (Figure 17b) was measured as the uptake of $^{35}\text{SO}_4^{2-}$ into macromolecular material as described (Taskiran, et al., 1994, Biophys. Res. Commun. 200:142-148). Results are shown in each case as percent of control. Values given are means \pm S.E. Numbers of knees are shown above each column.

Figure 18 shows effects of IRAP gene transfer on leucocytosis. Experiment conditions were identical to those described in Figure 15. Numbers of leukocytes in the lavage fluids were determined with a hemocytometer. Values shown are means \pm S.E. Numbers of knees are shown above each column. Asterisks denote values which differ at $p < 0.05$ (t-test).

DETAILED DESCRIPTION OF THE INVENTION

As used herein, the term "patient" includes members of the animal kingdom including but not limited to human beings.

As used herein, the term "mammalian host" includes members of the animal kingdom including but not limited to human beings.

As used herein, the term "connective tissue" includes but is not limited to a ligament, a cartilage, a tendon, and a synovium of a mammalian host.

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As used herein, the term "DC-chol" means a cationic liposome containing cationic cholesterol derivatives. The "DC-chol" molecule includes a tertiary amino group, a medium length spacer arm (two atoms) and a carbamoyl linker bond as described in Biochem. Biophys. Res. Commun., 179:280-285 (1991), X. Gao and L. Huang.

5 As used herein, "SF-chol" is defined as a type of cationic liposome.

As used herein, the term "biologically active" used in relation to liposomes denotes the ability to introduce functional DNA and/or proteins into the target cell.

As used herein, the term "biologically active" in reference to a nucleic acid, protein, protein fragment or derivative thereof is defined as an ability of the nucleic acid or amino
10 acid sequence to mimic a known biological function elicited by the wild type form of the nucleic acid or protein.

As used herein, the term "maintenance", when used in the context of liposome delivery, denotes the ability of the introduced DNA to remain present in the cell. When
used in other contexts, it means the ability of targeted DNA to remain present in the targeted
15 cell or tissue so as to impart a therapeutic effect.

Connective tissues are difficult organs to target therapeutically. Intravenous and oral routes of drug delivery that are known in the art provide poor access to these connective tissues and have the disadvantage of exposing the mammalian host body systemically to the therapeutic agent. More specifically, known intra-articular injection of joints provides direct
20 access to a joint. However, most of the injected drugs have a short intra-articular half-life. The present invention solves these problems by introducing into the connective tissue of a mammalian host genes encoding for proteins that may be used to treat the mammalian host. More specifically, this invention provides a method for introducing into the connective tissue of a mammalian host genes encoding for proteins with anti-arthritic properties.

25 The present invention provides a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host which comprises employing recombinant techniques to produce a viral vector which contains the gene encoding for the product, and infecting the connective tissue cell of the mammalian host using the viral vector containing the gene coding for the product.

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This method preferably includes introducing the gene encoding the product into at least one cell of the connective tissue of the mammalian host for a therapeutic use.

One *ex vivo* method of treating a connective tissue disorder disclosed throughout this specification comprises initially generating a recombinant viral vector which contains a DNA
5 sequence encoding a protein or biologically active fragment thereof. This recombinant viral vector is then used to infect a population of *in vitro* cultured connective tissue cells, resulting in a population of transfected connective cells. These transfected connective tissue cell are then transplanted to connective tissue within a target joint space of a mammalian host, effecting subsequent expression of the protein or protein fragment within the joint space.

10 Expression of this DNA sequence of interest is useful in substantially reducing at least one deleterious joint pathology or indicia of inflammation normally associated with a connective tissue disorder.

The connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament, preferably synovial cells.

15 It is also preferred that a retroviral vector, such as MFG, be utilized as the viral vector.

Another preferred step in this *ex vivo* method is transplantation of transduced cells by by intraarticular injection.

It will be understood by the artisan of ordinary skill that the preferred source of cells
20 for treating a human patient are the patients own cells, such as autologous synovial cells.

In a preferred embodiment of this invention, the method as hereinbefore described includes employing as the gene a DNA sequence encoding a human interleukin-1 receptor antagonist protein (IRAP) or biologically active fragment thereof.

Therefore, a preferred embodiment of the present invention relates to a method of
25 using a DNA sequence encoding a biologically active interleukin-1 receptor antagonist (IRAP) or portion thereof for treatment of connective tissue joint pathologies. The DNA sequence encoding IRAP or a biologically active fragment thereof may be delivered to the connective tissue of a mammalian host by any combination of various vector strategies and transduction techniques disclosed throughout this specification.

A preferred method of the embodiment of delivering IRAP to a target joint space involves delivery of the IPAP gene to the synovial lining of a mammalian host through use of a retroviral vector with the *ex vivo* technique disclosed within this specification. In other words, a DNA sequence of interest encoding a functional IRAP protein or protein fragment is subcloned into a retroviral vector of choice, the recombinant viral vector is then grown to adequate titers and used to infect *in vitro* cultured synovial cells, and the transduced synovial cells, preferably autografted cells, are transplanted into the joint of interest, preferably by intra-articular injection.

In another embodiment of this invention, the method as hereinbefore described includes employing as the gene a Lac Z marker gene capable of encoding a beta-galactosidase.

In another embodiment of this invention, the method as hereinbefore described includes employing as the gene a gene capable of encoding a soluble interleukin-1 receptor.

In another embodiment of this invention, the method as hereinbefore described includes employing as the gene a gene capable of encoding a soluble TNF- α receptor.

Another embodiment of this invention includes the method as hereinbefore described including employing as the gene a gene capable of encoding at least proteinase inhibitor. More specifically, this method preferably includes employing a tissue inhibitor of a metalloproteinases as the proteinase inhibitor.

Another embodiment of this invention includes the method as hereinbefore described including employing as the gene a gene capable of encoding at least one cytokine. More specifically, this method includes employing as the cytokine at least one material selected from the group consisting of interleukin-1 alpha (IL-1 α), interleukin-1 beta (IL-1 β), interleukin-2 (IL-2), interleukin-3 (IL-3), interleukin-4 (IL-4), interleukin-5 (IL-5), interleukin-6 (IL-6), interleukin-7 (IL-7), interleukin-8 (IL-8), interleukin-9 (IL-9), interleukin-10 (IL-10), interleukin-11 (IL-11), interleukin-12 (IL-12), tumor necrosis factor- α (TNF- α), and tumor necrosis factor- β (TNF- β).

A further embodiment of this invention includes a method as hereinbefore described including employing as the cytokine at least one transforming growth factor. More specifically, this method includes employing as the transforming growth factor a growth

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factor selected from the group consisting of TGF-beta₁, TGF-beta₂, TGF-beta₃, and TGF-alpha. Each transforming growth factor is commercially available from R & D Systems, 614 McKinley Place, N.E., Minneapolis, MN 55413.

5 In another embodiment of this invention, the method as hereinbefore described includes employing as the cytokine at least one fibroblast growth factor. The fibroblast growth factors are also commercially available from R & D Systems, 614 McKinley Place, N.E., Minneapolis, MN 55413.

10 Another embodiment of this invention includes the method as hereinbefore described including employing as the viral vector a retroviral vector. More specifically, this method includes employing as the retroviral vector at least one material selected from the group consisting of MFG and BAG. A preferred embodiment of this invention includes providing the method as hereinbefore described including employing as the gene a gene capable of encoding a human interleukin-1 receptor antagonist protein and employing MFG as the retroviral vector.

15 Another preferred embodiment of this invention includes the method as hereinbefore described including employing a Lac Z marker gene as the gene capable of encoding a beta-galactosidase and employing MFG as the retroviral vector.

20 Another preferred embodiment of this invention provides the method as hereinbefore described including employing a Lac Z neo marker gene as the gene capable of encoding a beta-galactosidase and employing BAG as the retroviral vector.

In a most preferred embodiment of this invention, the method as hereinbefore described includes employing a retroviral vector selected from the group consisting of MFG and BAG and includes employing as the gene a gene capable of encoding a soluble interleukin-1 receptor.

25 In another embodiment of this invention, a method as hereinbefore described is provided including employing as the gene a gene capable of encoding at least one proteinase inhibitor and including employing as the retroviral vector at least one material selected from the group consisting of MFG and BAG.

30 In another embodiment of this invention, a method as hereinbefore described is provided which includes employing as the retroviral vector at least one material selected

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from the group consisting of MFG and BAG and including employing as the gene a gene capable of encoding at least one cytokine as hereinbefore described.

5 In another embodiment of this invention, a method is provided for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host which comprises employing recombinant techniques to produce a viral vector which contains the gene encoding for the product and infecting the connective tissue cell of the mammalian host using the viral vector containing the gene coding for the product, wherein the viral vector is at least one vector selected from the group consisting of an adeno-associated virus, an adenovirus, and a herpes virus, such as herpes
10 simplex type-1 or herpes simplex type-2. This method includes employing as the gene a gene capable of encoding at least one material selected from the group which includes (a) a human interleukin-1 receptor antagonist protein, (b) a soluble interleukin-1 receptor, (c) a Lac Z marker gene capable of encoding a beta-galactosidase, (d) at least one proteinase inhibitor and (e) at least one cytokine. More specifically, this method includes employing a
15 tissue inhibitor of metalloproteinases as the proteinase inhibitor and includes employing as the cytokine at least one of the materials selected from the group which includes (a) at least one transforming growth factor selected from the group consisting of TGF-beta₁, TGF-beta₂, TGF-beta₃, and TGF-alpha, (b) at least one fibroblast growth factor, (c) IL-1 α , (d) IL-1 β , (e) IL-2 (f) IL-3, (g) IL-4, (h) IL-5, (i) IL-6, (j) IL-7, (k) IL-8, (l) IL-9, (m) IL-10, (n) IL-11,
20 (o) IL-12, (p) TNF- α , and (q) TNF- β

Another embodiment of this invention includes the method as hereinbefore described including introducing the gene into a connective tissue which tissue is selected from the group consisting of a ligament, a cartilage, a tendon, and a synovium. It is preferable that this method includes employing a cruciate ligament as the ligament. Most preferable this
25 method includes employing as the cruciate ligament a ligament selected from the group consisting of an anterior cruciate ligament and a posterior cruciate ligament.

Another embodiment of this invention includes the method as hereinbefore described including employing as the gene a gene having DNA that is capable of maintenance and expression.

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A further embodiment of this invention includes the method as hereinbefore described including introducing the gene into the cell *in vitro*. This method includes subsequently transplanting the infected cell into the mammalian host. This method also includes after effecting the infecting of the connective tissue cell but before the transplanting of the infected cell into the mammalian host, storing the infected connective tissue cell. It will be appreciated by those skilled in the art that the infected connective tissue cell may be stored frozen in 10 percent DMSO in liquid nitrogen. This method includes employing a method to substantially prevent the development of arthritis in a mammalian host having a high susceptibility of developing arthritis.

The method of this invention includes employing the method on an arthritic mammalian host for a therapeutic use. This method includes employing a method to repair and regenerate the connective tissue which tissue is selected from the group consisting of a ligament, a cartilage, a tendon, and a synovium. This method includes employing the method on a mammalian host that is a human being.

Another embodiment of this invention includes a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host as hereinbefore described including effecting *in vivo* the infection of the cell by introducing the viral vector containing the gene coding for the product directly into the mammalian host. Preferably, this method includes effecting the direct introduction into the mammalian host by intra-articular injection. This method includes employing the method to substantially prevent a development of arthritis in a mammalian host having a high susceptibility of developing arthritis. This method also includes employing the method on an arthritic mammalian host for therapeutic use. Further this method as includes employing the method to repair and regenerate the connective tissue as hereinbefore defined.

In yet another embodiment of this invention, a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host includes employing non-viral means for introducing the gene encoding for the product into the connective tissue cell. This method includes employing non-viral means selected from the group consisting of at least one liposome,

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$\text{Ca}_3(\text{PO}_4)_2$, electroporation, and DEAE-dextran. This method includes employing as the liposome a material selected from the group consisting of DC-chol and SF-chol.

It will be understood that the method of this invention of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host that includes employing non-viral means for introducing the gene encoding for the product into the connective tissue cell is a non-infectious delivery system. An advantage of the use of a non-infectious delivery system is the elimination of insertional mutagenesis and virally induced disease.

It will be appreciated by those skilled in the art, that the viral vectors employing a liposome are not limited by cell division as is required for the retroviruses to effect infection and integration of connective tissue cells. This method employing non-viral means as hereinbefore described includes employing as the gene a gene capable of encoding at least one of the following materials selected from the group which includes (a) a human interleukin-1 receptor antagonist protein, (b) a Lac Z marker gene capable of encoding a beta-galactosidase, (c) a soluble interleukin-1 receptor, (d) at least one proteinase inhibitor, (e) at least one transforming growth factor, and (f) at least one cytokine. More specifically, this method includes employing as the cytokine a cytokine selected from the group which includes IL-1 α , IL-1 β , IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, TNF- α , and TNF- β , and at least one fibroblast growth factor. Preferably, IL-4 and IL-10 are the selected cytokines. Preferably, this method includes employing as the transforming growth factor a growth factor selected from the group consisting of TGF-beta₁, TGF-beta₂, TGF-beta₃, and TGF-alpha.

High levels of collagenase and other tissue metalloproteinases, such as stromelysin and gelatinase can be expressed in the presence of IL-1 within connective tissue.

Collagenase, stromelysin, and gelatinase are inhibited by the protein TIMP ("Tissue Inhibitor of MetalloProteinases"). Therefore, another preferred embodiment of this invention includes providing the method employing non-viral means as hereinbefore described which includes employing a tissue inhibitor of metalloproteinases (TIMP) as the proteinase inhibitor. This method employing non-viral means for introducing the gene encoding for the product into the connective tissue cell as hereinbefore described includes introducing the gene into the

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connective tissue which tissue is selected from the group consisting of a ligament, a cartilage, a tendon, and a synovium. Preferably, this method includes employing a cruciate ligament as the ligament. The cruciate ligament is selected from the group consisting of an anterior cruciate ligament and an posterior cruciate ligament. Of course, a gene encoding a TIMP
5 protein or biologically active fragment thereof could be delivered to the target connective tissue by any combination of means disclosed in this specification.

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various antiadhesion molecules so as to
10 inhibit cell-cell and cell-matrix interactions. Examples of such proteins or protein fragments include but are not limited to soluble ICAM-1 and soluble CD44.

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various cartilage growth factors,
15 including but not limited to IGF-1 and TGF- β .

Another embodiment of the present invention is delivery of a DNA sequence of interest to the connective tissue of a mammalian host by any of the methods disclosed within this specification so as to effect *in vivo* expression of various free radical antagonists, thus preventing the deleterious effects of free radical formation within the afflicted joint.
20 Examples include but are not limited to the superoxide dismutase and proteins or protein fragments which inhibit NO.

A further embodiment of this invention provides for an animal model to study connective tissue pathologies and indices of systemic inflammation. This model utilizes either *ex vivo* or *in vivo* delivery of at least one gene or DNA sequence of interest encoding a
25 product into a least one cell of a connective tissue of a mammalian host. Examples of joint pathologies which can be studied in the present invention include, but are by no means limited to, joint pathologies such as leukocytosis, synovitis, cartilage breakdown and suppression of cartilage matrix synthesis. Examples of indices of systemic inflammation which include, but are by no means limited to, erythrocyte sedimentation rate, fever and
30 weight loss.

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A particular embodiment of the present invention which relates to such an animal model is utilization of the *ex vivo* based delivery of a DNA sequence encoding human IL-1 β gene to the synovial lining of the rabbit knee. In this embodiment, the human IL-1 β gene was subcloned into the MFG retroviral vector by known methods, resulting in MFG-IL-1 β .
5 This recombinant retroviral construct was used to transduce autologous synovial cells cultured *in vitro*. These transduced cells were then delivered to the rabbit knees as described throughout this specification. Delivery of the human IL-1 β gene to the synovial lining of the rabbit knee in this fashion caused a severe, chronic, monoarticular arthritis. Pathologies included leukocytosis, synovitis, cartilage breakdown and suppression of cartilage matrix
10 synthesis. Various systemic indices of inflammation were also effected, including an increased erythrocyte sedimentation rate, fever and weight loss.

In another example of this particular embodiment of the present invention, the human IL-1 β gene was subcloned into a DNA plasmid vector, downstream of a CMV promoter. This CMV-IL-1 β plasmid construct was encapsulated in liposomes and delivered
15 to a target joint space as described in Example X. Forty eight hours subsequent to injection 1 ng of IL-1 β was recovered from the knee joint area.

An animal model as described and exemplified in this specification measures the ability of various gene therapy applications disclosed throughout this specification to withstand challenges from known causative agents (such as IL-1 β) of joint pathologies and
20 inflammatory side effects.

Another preferred method of the present invention involves direct *in vivo* delivery of the IRAP gene to the synovial lining of a mammalian host through use of either an adenovirus vector, adeno-associated virus (AAV) vector or herpes-simples virus (HSV) vector. In other words, a DNA sequence of interest encoding a functional IPAP protein or
25 protein fragment is subcloned into the respective viral vector, the IRAP containing viral vector is then grown to adequate titers and directed into the joint space, preferably by intra-articular injection. A retroviral-IRAP construct, such as MFG-IRAP may also be utilized to directly target previously inflamed connective tissue cells within the joint space.

Direct intra-articular injection of a DNA molecule containing the gene of interest into
30 the joint results in transfection of the recipient synovial cells and hence bypasses the

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requirement of removal, *in vitro* culturing, transfection, selection, as well as transplanting the DNA vector containing - synoviocytes (as disclosed in the Example section) to promote stable expression of the heterologous gene of interest. Methods of presenting the DNA molecule to the target connective tissue of the joint includes, but is not limited to, encapsulation of the DNA molecule into cationic liposomes, subcloning the DNA sequence of interest in a retroviral vector as described throughout this specification, or the direct injection of the DNA molecule itself into the joint. The DNA molecule, regardless of the form of presentation to the knee joint, is preferably presented as a DNA vector molecule, either as recombinant viral DNA vector molecule or a recombinant DNA plasmid vector molecule. Expression of the heterologous gene of interest is ensured by inserting a promoter fragment active in eukaryotic cells directly upstream of the coding region of the heterologous gene. One of ordinary skill in the art may utilize known strategies and techniques of vector construction to ensure appropriate levels of expression subsequent to entry of the DNA molecule into the synovial tissue. *In vivo* delivery of various viral and non-viral vectors to the rabbit knee joint are described in Example XV.

Another embodiment of this invention provides the method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host which includes employing non-viral means as hereinbefore described and includes employing as the gene a gene having DNA that is capable of maintenance and expression.

In yet a further embodiment of this invention, the method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host is provided that includes employing non-viral means for introducing the gene encoding for the product into the connective tissue cell *in vitro* and includes subsequently transplanting the cell having the gene into the mammalian host.

Another embodiment of this invention provides a method including after introducing the gene encoding for the product in the connective tissue cell and before the transplanting of the connective tissue cell having the gene into the mammalian host, storing the connective tissue cell having the gene. This method includes storing connective tissue cell frozen in 10 percent DMSO in liquid nitrogen. This method includes employing a method to substantially

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prevent the development of arthritis in a mammalian host having a high susceptibility of developing arthritis. Further, this method includes employing the method on an arthritic mammalian host for a therapeutic use. This method includes employing the method to repair and regenerate the connective tissue which tissue is selected from the group consisting of a ligament, a cartilage, a tendon, and a synovium.

A further embodiment of this invention provides a method of introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host which includes employing non-viral means *in vivo* for directly introducing the gene encoding for the product into the connective tissue cell of the mammalian host. The non-viral means is selected from the group consisting of at least one liposome, $\text{Ca}_3(\text{PO}_4)_2$ and DEAE-dextran. Preferably, this method includes effecting the *in vivo* introduction into the mammalian host by intra-articular injection. This method includes employing the method to substantially prevent the development of arthritis in a mammalian host having a high susceptibility of developing arthritis. Further, this method includes employing the method on an arthritic mammalian host for a therapeutic use. This method also includes employing the method to repair and regenerate the connective tissue which tissue is selected from the group consisting of a ligament, a cartilage, a tendon, and a synovium.

Another embodiment of the present invention is a method to produce an animal model for the study of connective tissue pathology. As will be understood by those skilled in the art, over-expression of interleukin-1 in the joint of a mammalian host is generally responsible for the induction of an arthritic condition. This invention provides a method for producing an animal model using the hereinbefore described gene transfer technology of this invention. Preferably, the method of this invention provides a method for producing an animal model using the hereinbefore described gene transfer technology of this invention to effect an animal model for arthritis. For example, constitutive expression of interleukin-1 in the joint of a rabbit following the method of gene transfer provided for by this invention leads to the onset of an arthritic condition. It will be appreciated by those skilled in the art that this rabbit model is suitable for use for the testing of therapeutic agents. This method includes introducing at least one gene encoding a product into at least one cell of a connective tissue

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of a mammalian host comprising (a) employing recombinant techniques to produce a viral vector which contains the gene encoding for the product and (b) infecting the connective tissue cell of the mammalian host using the viral vector containing the gene coding for the product for effecting the animal model. This method includes employing as the gene a material selected from the group consisting of a cytokine and a proteinase. This method includes employing as the cytokine a material selected from the group consisting of IL-1 α , IL-1 β and TNF- α . This method includes employing as the proteinase a matrix metalloproteinase. The matrix metalloproteinase is an enzyme selected from the group consisting of a collagenase, a gelatinase and a stromelysin. It will be apparent that use of the term "a collagenase, a gelatinase and a stromolysin" is meant to include the plural, and not be limited to the singular. It is well known in the art that numerous collagenases, gelatinases and stromolysins could be employed as a matrix metalloproteinase in the present invention. A further embodiment of this invention provides a method to produce an animal model for the study of connective tissue pathology which includes employing non-viral means for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for effecting the animal model. The non-viral means is selected from the group consisting of at least one liposome, Ca₃(PO₄)₂, electroporation, and DEAE-dextran. This method includes employing as the gene a material selected from the group consisting of a cytokine and a proteinase. This method includes employing as the cytokine a material selected from the group consisting of IL-1 α , IL-1 β and TNF- α . This method also includes employing as the proteinase a matrix metalloproteinase. The matrix metalloproteinase includes an enzyme selected from at least one of the group consisting of a collagenase, a gelatinase, and a stromelysin.

A further embodiment of the present invention includes employing as the gene a gene capable of encoding at least one of the materials which is selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or a biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or a biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or a biologically active derivative or fragment thereof, (d) a proteinase inhibitor, (e) a soluble tumor necrosis factor receptor protein or a biologically

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active derivative or fragment thereof and (f) a cytokine, and employing as the DNA vector any DNA vector, preferably a plasmid or viral vector, known to one of ordinary skill in the art capable of stable maintenance within the targeted cell or tissue upon delivery, regardless of the method of delivery utilized. In one embodiment of the invention, synoviocytes are

5 transfected *in vivo* subsequent to direct intra-articular injection of a DNA molecule containing the gene of interest into the joint. Transfection of the recipient synovial cells bypasses the requirement of removal, culturing, *in vitro* transfection, selection and transplanting the DNA vector containing - synoviocytes (as disclosed in the Example section) to promote stable

10 expression of the heterologous gene of interest. Methods of injecting the DNA molecule into the joint includes, but is not limited to, encapsulation of the DNA molecule into cationic liposomes or the direct injection of the DNA molecule itself into the joint. Expression of the heterologous gene of interest subsequent to *in vivo* transfection of the synovial tissue is ensured by inserting a promoter fragment active in eukaryotic cells directly upstream of the coding region of the heterologous gene. One of ordinary skill in the art may utilize known

15 strategies and techniques of vector construction to ensure appropriate levels of expression subsequent to entry of the DNA molecule into the synovial tissue. As an example, and not a limitation, of the present invention, a DNA plasmid vector containing the IL-1 β coding sequence ligated downstream of the CMV promoter was encapsulated within liposomes and injected into the knee joints of recipient rabbits. Interleukin-1 beta was expressed in synovial

20 tissue, as significant amounts of interleukin-1 beta was recovered from the synovial tissue within the region of intra-articular injection.

A further embodiment of this invention provides an additional method for introducing at least one gene encoding a product into at least one cell of a connective tissue of a mammalian host for use in treating the mammalian host. This additional method includes

25 employing the biologic means of utilizing a virus to deliver the DNA vector molecule to the target cell or tissue. Preferably, the virus is a psuedovirus, the genome having been altered such that the psuedovirus is capable only of delivery and stable maintenance within the target cell; but not retaining an ability to replicate within the target cell or tissue. The altered viral genome is further manipulated by recombinant DNA techniques such that the viral genome

30 acts as a DNA vector molecule which contains the heterologous gene of interest to be

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expressed within the target cell or tissue. This method also includes employing as the gene a gene capable of encoding at least one of the materials selected from the group which includes (a) a human interleukin-1 receptor antagonist protein or biologically active derivative or fragment thereof, (b) a Lac Z marker gene capable of encoding a beta-galactosidase protein or biologically active derivative or fragment thereof, (c) a soluble interleukin-1 receptor protein or biologically active derivative or fragment thereof, (d) a proteinase inhibitor and (e) a soluble tumor necrosis factor receptor protein or a biologically active derivative or fragment thereof and (f) a cytokine.

The following examples are offered by way of illustration of the present invention, and not by way of limitation.

EXAMPLE I

Packaging of AAV

The only cis-acting sequences required for replication and packaging of recombinant adeno-associated virus (AAV) vector are the AAV terminal repeats. Up to 4 kb of DNA can be inserted between the terminal repeats without effecting viral replication or packaging. The virus rep proteins and viral capsid proteins are required in trans for virus replication as is an adeno-associated virus helper. To package a recombinant AAV vector, the plasmid containing the terminal repeats and the therapeutic gene is co-transfected into cells with a plasmid that expresses the rep and capsid proteins. The transfected cells are then infected with adeno-associated virus and virus isolated from the cells about 48-72 hours post-transfection. The supernatants are heated to about 56° Centigrade to inactivate the adeno-associated virus, leaving a pure virus stock of recombinant AAV.

EXAMPLE II

Electroporation

The connective tissue cells to be electroporated are placed into Hepes buffer saline (HBS) at a concentration of about 10^7 cells per ml. The DNA to be electroporated is added at a concentration of about 5-20 ug/ml of HBS. The mixture is placed into a cuvette and inserted into the cuvette holder that accompanies the Bio-RAD electroporation device (1414 Harbour Way South, Richmond, CA 94804). A range between about 250 and 300 volts at a capacitance of about 960 ufarads is required for introduction of DNA into most eukaryotic

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cell types. Once the DNA and the cells are inserted into the Bio-RAD holder, a button is pushed and the set voltage is delivered to the cell-DNA solution. The cells are removed from the cuvette and replated on plastic dishes.

EXAMPLE III

5 The cDNA encoding the human interleukin-1 receptor antagonist (IRAP) was inserted into the NcoI and BamHI cloning sites of the retroviral vector MFG as shown in Figure 1. Specifically, a PstI to BamHI fragment from the IRAP cDNA was linked to a synthetic oligonucleotide adapter from the NcoI site (representing the start site of translation for IRAP) to the PstI site (approximately 12 base pairs downstream from the NcoI site) to the MFG backbone digested at NcoI and BamHI in a three part ligation reaction. This three part
10 ligation involving a synthetic oligo and two DNA fragments is well known by those skilled in the art of cloning. LTR means long terminal repeats, 5'SD means 5' splice donor, 3'SA means 3' splice acceptor. The straight arrow and the crooked arrow in Figure 1 represent unspliced and spliced messenger RNAs respectively. IRAP is encoded by the spliced
15 message.

Figure 2 shows the cDNA encoding the human interleukin-1 receptor antagonist protein (IRAP) with a selectable neo gene marker. Figure 3 shows a low power micrograph of synovium recovered from the knee of a rabbit one month after intra-articular injection of Lac Z⁺, neo⁺ synoviocytes. Tissue was stained histochemically for the presence of
20 beta-galactosidase. This micrograph counterstained with eosin revealed an area of intensely stained, transplanted cells demonstrating that these cells have colonized the synovial lining of the recipient joint.

EXAMPLE IV

Animal Models

25 The methods of this invention of transferring genes to the synovia of mammalian joints permit the production and analysis of joint pathologies that were not previously possible. This is because the only other way of delivering potentially arthritogenic compounds to the joint is by intra-articular injection. Not only are such compounds quickly cleared from joints, but the effects of bolus injections of these compounds do not accurately
30 mimic physiological conditions where they are constantly produced over a long period of

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time. In contrast, the gene transfer technologies of this invention permit selected proteins of known or suspected involvement in the arthritic process to be expressed intra-articularly over an extended period of time, such as for example, at least a three month period. The animal models of this invention therefore permits the importance of each gene product to the arthritic process to be evaluated individually. Candidate genes include, but are not restricted to, those coding for cytokines such as interleukin-1 (IL-1) alpha, IL-1 beta, and TNF-alpha, and matrix metalloproteinases such as collagenases, gelatinases and stromelysins.

Additionally, the gene transfer techniques of this invention are suitable for use in the screening of potentially therapeutic proteins. In this use, the animal models of the invention are initiated in joints whose synovia express gene coding for potential anti-arthritic proteins. Candidate proteins include, but are not restricted to, inhibitors of proteinases such as, for example, the tissue inhibitor of metalloproteinases, and cytokines such as, for example, transforming growth factor-beta.

EXAMPLE V

Method For Using Synoviocytes As A Delivery System For Gene Therapy

Rabbits are killed by intravenous injection of 4 ml nembutol, and their knees quickly shaved. Synovia are surgically removed from each knee under aseptic conditions, and the cells removed from their surrounding matrix by sequential digestion with trypsin and collagenase (0.2% w/v in Gey's Balanced Salt Solution) for about 30 minutes and about 2 hours, respectively. The cells recovered in this way are seeded into 25 cm² culture flasks with about 4 ml of Ham's F₁₂ nutrient medium supplemented with 10% fetal bovine serum, about 100 U/ml penicillin and about 100 µg/ml streptomycin, and incubated at about 37° in an atmosphere of 95% air, 5% CO₂. Following about 3-4 days incubation, the cells attain confluence. At this stage, the culture medium is removed and the cell sheet washed twice with approximately 5 mls of Grey's Balanced Salt Solution to remove non-adherent cells such as lymphocytes. The adherent cells are then treated with trypsin (0.25% w/v in balanced salt solution). This treatment detaches the fibroblastic, Type B synoviocytes, but leaves macrophages, polymorphonuclear leukocytes and the Type A synoviocytes attached to the

culture vessel. The detached cells are recovered, re-seeded into 25 cm² culture vessels at a 1:2 split ratio, medium is added and the culture returned to the incubator. At confluence this procedure is repeated.

5 After the third such passage, the cells are uniformly fibroblastic and comprise a homogeneous population of Type B synoviocytes. At this stage, cells are infected with the retroviral vector.

10 Following infection, cells are transferred to fresh nutrient medium supplemented with about 1 mg/ml G418 (GIBCO/BRL, P.O. Box 68, Grand Island, NY 14072-0068) and returned to the incubator. Medium is changed every three days as neo⁻ cells die and the neo⁺ cells proliferate and attain confluency. When confluent, the cells are trypsinized and subcultured as described above. One flask is set aside for staining with X-gal to confirm that the neo⁺ cells are also Lac Z⁺. When the subcultures are confluent, the medium is recovered and tested for the presence of IRAP, soluble IL-1R or other appropriate gene products as hereinbefore described. Producing synoviocyte cultures are then ready for
15 transplantation.

The day before transplantation, the cells are recovered by trypsinizing, as hereinbefore described. These cells are then suspended in nutrient medium, and incubated overnight in an untreated plastic centrifuge tube. Under these conditions, the cells do not adhere, but they regenerate their cell surface proteins that were removed by trypsinizing.

20 The following morning, the cells are recovered by centrifuging, washed several times by resuspension in Gey's Balanced Salt Solution and finally resuspended at a concentration of about 10⁶-10⁷ cells/ml in Grey's solution. Approximately 1 ml of this suspension is then introduced into the knee joint of a recipient rabbit by intra-articular injection. For this purpose a 1 ml syringe with a 25-gauge hypodermic needle is used. Injection is carried out
25 through the patellar tendon. Experiments in which radiopaque dye was injected have confirmed that this method successfully introduces material into all parts of the joint.

Variations on the disclosed harvesting, culture and transplantation conditions in regard to the numerous examples presented within this specification will be evident upon inspection of this specification. Several tangential points may be useful to one practicing the ex vivo
30 based gene therapy portion of the disclosed invention:

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(1) If the yield of synoviocytes from the harvested synovial tissue is poor, the surgical technique may be at fault. The synovium has a strong tendency to retract when cut. Therefore, the inner capsule is grasped firmly, and with it the synovium, while excising this tissue. A small (about 2 mm) transverse incision can be made inferiorly, followed by sliding one point of the forceps into the joint space so that the synovium and inner capsule are sandwiched between the points of the forceps. The tissue is then excised without releasing the tissue thus preventing retraction of the synovium.

(2) A two compartment digestion chamber may be used to initially separate the cells from extracellular debris. In lieu of this choice, synovial tissue may be digested in a single chamber vessel and filtered through a nylon monofilament mesh of 45 μm pore size.

(3) When resuspending cells, the smallest amount of medium possible can be used to prevent formation of clumps of cells, which are difficult to separate once formed.

(4) During trypsinization, synoviocytes can lose the stellate/dendritic morphology that they possess in adherence, and assume a rounded shape. The cells initially will detach in clumps of rounded cells; one may allow the majority of cells to separate from each other before stopping trypsinization.

(5) Synoviocytes may be transduced with multiple transgenes by use of retroviral vectors containing multiple transgenes or by sequential transduction by multiple retroviral vectors. In sequential transduction, the second transduction should be made following selection, when applicable, and passage after the first transduction.

(6) As the synovium is a well-innervated structure, intra-articular injection can be painful, especially if done rapidly. Intra-articular injection of a 1 ml volume should take 10 to 15 seconds.

(7) In the animal model, the depth of the needle stick should not exceed 1 cm during intraarticular injection, and depression of the syringe plunger should meet with little to no resistance. Resistance to advancement of the syringe plunger indicates that the tip of the needle is not in the joint space.

(8) In the animal model, to retrieve a useful volume of the injected Gey's solution during joint lavage, the needle should not be inserted too deeply, otherwise it may penetrate the posterior capsule and may lacerate the popliteal artery. Firm massage of the

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suprapatellar, infrapatellar, and lateral aspects of the knee during aspiration helps to increase the amount of fluid recovered; in general, it should be possible to recover ≥ 0.5 ml of fluid. When knees are badly inflamed, lavage is often difficult because of the presence of large numbers of leukocytes, fibrin, and other debris in the joint. Under such conditions the only recourse is to sacrifice the animal and recover the Gey's solution surgically.

EXAMPLE VI

The method of Example V for producing generally uniformly fibroblastic cells of a homogeneous population of Type B synoviocytes was followed to effect growing cultures of lapine synovial fibroblasts. These growing cultures of lapine synovial fibroblasts were subsequently infected with an amphotropic retroviral vector carrying marker genes coding for beta-galactosidase (Lac Z) and resistance to the neomycin analogue G418 (neo⁺). Following infection and growth in selective medium containing about 1 mg/ml G418, all cells stained positively in a histochemical stain for beta-galactosidase.

Neo selected cells carrying the Lac Z marker gene were transplanted back into the knees of recipient rabbits to examine the persistence and expression of these genes *in vivo*. Two weeks following transplantation, islands of Lac Z⁺ cells within the synovium of recipient knees were observed. This confirmed the ability of the method of this invention to introduce marker genes into rabbit synovia and to express them *in situ*.

EXAMPLE VII

Neo-selected, Lac Z⁺ synoviocytes were recovered from cell culture, suspended in Gey's Balanced Salt Solution and injected intra-articularly into the knee joints of recipient rabbits (about 10^5 - 10^7 cells per knee). Contralateral control knees received only a carrier solution. At intervals up to 3 months following transplant, the rabbits were killed and their synovia and surrounding capsule recovered. Each sample may be analyzed in three ways. A third of the synovium was stained histochemically *en masse* for the presence of beta-galactosidase. A second portion may be used for immunocytochemistry using antibodies specific for bacterial beta-galactosidase. The final portion may be digested with trypsin and collagenase, and the cells thus recovered cultured in the presence of G418.

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Staining of the bulk synovial tissue revealed extensive areas of Lac Z⁺ cells, visible to the naked eye. Control synovia remained colorless. Histochemical examination of synovia revealed the presence of islands of cells staining intensely positive for beta-galactosidase. These cells were present on the superficial layer of the synovial lining, and were absent from control synovia. From such tissue it was possible to grow Lac Z⁺, neo⁺ cells. Cells recovered from control tissue were Lac Z⁻ and died when G418 was added to the culture. This indicates that the transplanted, transduced synovial fibroblasts have successfully recolonized the synovia of recipient joints, and continue to express the two marker genes, Lac Z and neo. Maintaining intra-articular Lac Z and neo expression in transplanted synoviocytes has been effected for 3 months using primary cells and one month using the HIG-82 cell line.

EXAMPLE VIII

Based upon the methods of the hereinbefore presented examples, and employing standard recombinant techniques well known by those skilled in the art, the human IRAP gene was incorporated into an MFG vector as shown in Figure 1. Following the infection of synoviocyte cultures of rabbit origin with this viral vector, IRAP was secreted into the culture medium.

Western blotting, well known by those skilled in the art, was carried out using an IRAP-specific rabbit polyclonal antibody that does not recognize human or rabbit IL-1 alpha or IL-1 beta, or rabbit IRAP. Figure 4 shows a Western blot which sets forth the production of IRAP by four cultures of HIG-82 cells infected with MFG-IRAP. Three forms of the IRAP are present: a non-glycosylated form which runs with recombinant standards, and two larger glycosylated forms. The results of the Western blotting shown in Figure 4 demonstrated that IRAP was produced by HIG-82 synoviocyte cell line (Georgescu, 1988, In Vitro 24: 1015-1022) following infection with the MFG-IRAP vector of this invention. The Western blotting of Figure 4 shows the IRAP concentration of the conditioned medium is as high as 50 ng/ml. This is approximately equal to 500 ng IRAP/10⁶ cells/day. Lane 1 and Lane 2 of Figure 4 show that the recipient synovia tissue secrete substantial amounts of HIG-IRAP at 3 days (Lane 2) and 6 days (Lane 1). Lane 3 shows human recombinant IRAP. Lane 6 indicates that rabbit synovial cells produce a larger glycosylated version of

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this molecule after infection with MFG-IRAP. Lane 7 indicates that native rabbit synovial cells do not produce this glycosylated form.

Figure 5 shows that medium conditioned by IRAP⁺ synoviocytes blocks the induction of neutral metalloproteinases in articular chondrocytes exposed to recombinant human IL-1 beta. Chondrocytes normally secrete 1 U/10⁶ cells, or less, gelatinase into their culture media. Figure 5 shows that when to about 5 U/ml or 10 U/ml IL-1 are added, gelatinase production increases to over 4 U and 6U/10⁸ cells, respectively. Addition of medium conditioned by MFG-IRAP-infected HIG-82 cells employed by the method of this invention suppressed gelatinase production by IL-1 treated chondrocytes. With 5 U/ml IL-1 (Figure 5, right panel) inhibition was 100% for one culture and 41% for the other. With 10 U/ml IL-1, inhibition was reduced to 38% and 18% (Figure 5, left panel) as is expected of a competitive inhibitor. These data demonstrate that the IRAP produced by HIG-82 cells infected with MFG-IRAP is biologically active.

EXAMPLE IX

This example demonstrates the uptake and expression of Lac Z gene by synoviocytes using infection by a liposome (lipofection). A six well plate containing synovioyte cultures were transduced with the Lac Z gene by lipofection. The content of each well is as follows:

Well 1	Control cells, treated with liposomes alone
Well 2	Control cells, treated with DNA alone
Well 3	DNA + 150 nmole liposomes
Well 4	DNA + 240 nmole liposomes
Well 5	DNA + 300 nmole liposomes
Well 6	DNA + 600 nmole liposomes

Wells 3-6 containing sub-confluent cultures of synovial fibroblasts were infected with 6 ug of DNA complexed with 150-600 nmoles/well of "DC-chol" liposome or in the alternative, with "SF-chol". Three days later, cells were stained histochemically for expression of beta-galactosidase (Figure 6).

Table 1 shows the results of using the liposomes "DC-chol" and "SF-chol" in converting synovioyte cultures to the Lac Z⁺ phenotype without selection. Table 1 sets forth that the "DC-chol" liposome in a concentration of about 300 nmole/well converted

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generally 30% of the synovial cells in synoviocyte cultures to the Lac Z⁺ phenotype without selection. Reduced expression was shown in Well 6 for "DC-chol" due to the toxic effect of the high liposome concentration.

TABLE 1

% Lac Z⁺ Cells

	<u>Liposome, nmole/well</u>	<u>DC-chol</u>	<u>SF-chol</u>
5			
10	150	10	0.5
	240	22	1.0
	300	30	2.8
	600	NA	3.5

In another embodiment of this invention, a gene and method of using this gene provides for the neutralization of interleukin-1. Interleukin-1 is a key mediator of cartilage destruction in arthritis. Interleukin-1 also causes inflammation and is a very powerful inducer of bone resorption. Many of these effects result from the ability of interleukin-1 to increase enormously the cellular synthesis of prostaglandin E₂, the neutral proteinases -- collagenase, gelatinase, and stromelysin, and plasminogen activator. The catabolic effects of interleukin-1 upon cartilage are exacerbated by its ability to suppress the synthesis of the cartilaginous matrix by chondrocytes. Interleukin-1 is present at high concentrations in synovial fluids aspirated from arthritic joints and it has been demonstrated that intra-articular injection of recombinant interleukin-1 in animals causes cartilage breakdown and inflammation.

Interleukin-1 exists as several species, such as unglycosylated polypeptide of 17,000 Daltons. Two species have previously been cloned, interleukin-1 alpha and interleukin-1 beta. The alpha form has a pI of approximately 5, and the beta form has a pI around 7. Despite the existence of these isoforms, interleukin-1 alpha and interleukin-1 beta have substantially identical biological properties and share common cell surface receptors. The type I interleukin-1 receptor is a 80kDa (kilodalton) glycoprotein and contains an extracellular, interleukin-1 binding portion

of 319 amino acids which are arranged in three immunoglobulin-like domains held together by disulfide bridges as shown in Figure 7. A 21 amino acid trans-membrane domain joins the extracellular portion to the 217 amino acid cytoplasmic domain. Figures 8A-8C show the amino acid and nucleotide sequence of the human and mouse interleukin-1 receptors. In Figure 8B, the 21 amino acid trans-membrane region of the interleukin-1 receptor is marked by the thicker solid line. In Figures 8A and 8B, the position of the 5' and 3' oligonucleotides for PCR are marked by thinner short lines, respectively. The lysine amino acid just 5' to the trans-membrane domain to be mutated to a stop codon is marked by a solid circle in Figure 8B.

Synovium is by far the major, and perhaps the only, intra-articular source of interleukin-1 in the arthritic joint. Synovia recovered from arthritic joints secrete high levels of interleukin-1. Both the resident synoviocytes and infiltrating blood mononuclear cells within the synovial lining produce interleukin-1.

The present invention provides a method of using *in vivo* a gene coding for a truncated form of the interleukin-1 receptor which retains its ability to bind interleukin-1 with high affinity but which is released extracellularly and therefore inactive in signal transduction. The binding of this truncated and modified receptor to interleukin-1 inhibits the intra-articular activity of interleukin-1.

This method of using a gene encoding the extracellular interleukin-1 binding domain of an interleukin-1 receptor that is capable of binding to and neutralizing interleukin-1 includes employing a retroviral vector carrying a truncated interleukin-1 receptor gene which encodes a truncated and soluble active form of the receptor. The expression of the novel interleukin-1 receptor gene is controlled by regulatory sequences contained within the vector that are active in eukaryotic cells. This recombinant viral vector is transfected into cell lines stably expressing the viral proteins *in trans* required for production of infectious virus particles carrying the recombinant vector. These viral particles are used to deliver the recombinant interleukin-1 receptor to the recipient synovial cells by direct virus infection *in vivo*.

The soluble human interleukin-1 receptor to be inserted into the retroviral vector may be generated by a polymerase chain reaction (PCR). An oligonucleotide complementary to the 5' leader sequence of the human interleukin-1 receptor (GCGGATCCCCTCCTAGAAGCT; SEQ ID NO: 5) and an oligonucleotide complementary to a region just upstream from the

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transmembrane domain of the interleukin-1 receptor (GCGGATCCCATGTGCTACTGG; SEQ ID NO: 6) are used as primers for PCR. The primer for the region of the interleukin-1 receptor adjacent to the trans-membrane domain contains a single base change so that the lys codon at amino acid 336 (AAG) is changed to a stop codon (TAG). By inserting a translation stop codon just upstream from the transmembrane domain, a truncated form of interleukin-1 receptor that is secreted by the cell is generated. A BamHI recognition sequence (GGATCC) is added to the 5' end of the PCR primers, and following amplification, the resulting interleukin-1 receptor fragment is cloned into a BamHI site. A cDNA library from human T-cells is used as a source for the interleukin-1 receptor cDNA. To amplify the appropriate region of the interleukin-1 receptor from the cDNA library, the complementary primers are added to the DNA and 50 cycles of annealing, primer extension and denaturation are performed using a thermocycler and standard PCR reaction conditions well known by those persons skilled in the art. Following amplification of the interleukin-1 soluble receptor using the PCR process, the resulting fragment is digested with BamHI and inserted into the pLJ retroviral vector. The pLJ retroviral vector is available from A. J. Korman and R. C. Mulligan. See also Proc. Natl. Acad. Sci., Vol. 84, pp. 2150-2154 (April 1987) co-authored by Alan J. Korman, J. Daniel Frantz, Jack L. Strominger and Richard C. Mulligan. Restriction analysis was performed to determine the correct orientation of the insert.

The retrovirus vector carrying the truncated interleukin-1 receptor is transferred into the CRIP (Proc. Natl. Acad. Sci., Vol. 85, pp. 6460-6464 (1988), O. Danos and R. C. Mulligan) packaging cell line using a standard CaPO_4 transfection procedure and cells wherein the viral vector is stably integrated and is selected on the basis of resistance to the antibiotic G418. The viral vector containing the neomycin resistant (neo-r) gene is capable of imparting resistance of the cell line to G418. The CRIP cell line expresses the three viral proteins required for packaging the vector viral RNAs into infectious particles. Moreover, the viral particles produced by the CRIP cell line are able to efficiently infect a wide variety of mammalian cell types including human cells. All retroviral particles produced by this cell line are defective for replication but retain the ability to stably integrate into synovial cells thereby becoming an heritable trait of these cells. Virus stocks produced by this method are substantially free of contaminating helper-virus particles and are also non-pathogenic.

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More specifically, the truncated interleukin-1 gene can be inserted into a retroviral vector under the regulation of a suitable eukaryotic promoter such as the retroviral promoter already contained within the gene transfer vector, such as for example, the pLJ vector shown in Figure 9. Figure 9 shows the structure of the pLJ interleukin receptor retroviral vector and partial restriction endonuclease map. Reference numeral 10 shows the interleukin-1 receptor inserted into a retroviral vector. Reference numeral 12 indicates long terminal repeats (LTR's) at each end of the structure of the pLJ interleukin receptor retroviral vector shown in Figure 8. These LTR's regulate the viral transcription and expression of the interleukin-1 receptor. Bacterial gene encoding resistance to the antibiotic neomycin (neo-r) is shown at reference numeral 16. The Simian Virus 40 enhancer promoter (SV 40) is indicated at reference numeral 18, and regulates the expression of the neo-r gene. Reference numbers 20 and 22, respectively, show the sites wherein the resulting interleukin receptor fragment is cloned. It will be understood by those persons skilled in the art that other vectors containing different eukaryotic promoters may also be utilized to obtain a generally maximal level of interleukin-1 receptor expression. The vectors containing the truncated, and modified interleukin-1 receptor will be introduced into a retroviral packaging cell line (CRIP) by transfection and stable transformants isolated by selection for the expression of the neomycin resistance gene also carried by the pLJ vector. The CRIP cell line expresses all the proteins required for packaging of the exogenous retroviral RNA. Viral particles produced by the G418-selected CRIP cell lines will carry a recombinant retrovirus able to infect mammalian cells and stably express the interleukin-1 truncated receptor. The viral particles are used to infect synovial cells directly *in vivo* by injecting the virus into the joint space.

Another embodiment of this invention provides a method for using the hereinbefore described viral particles to infect in culture synovial cells obtained from the lining of the joint of a mammalian host. The advantage of the infection of synovial cells in culture is that infected cells harboring the interleukin-1 receptor retroviral construct can be selected using G418 for expression of the neomycin resistance gene. The infected synovial cells expressing the interleukin-1 receptor can then be transplanted back into the joint by intra-articular injection. The transplanted cells will express high levels of soluble interleukin-1 receptor in the joint space

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thereby binding to and neutralizing substantially all isoforms of interleukin-1, including interleukin-1 alpha and interleukin-1 beta.

The method used for transplantation of the synovial cells within the joint is a routine and relatively minor procedure used in the treatment of chronic inflammatory joint disease.

5 Although synovium can be recovered from the joint during open surgery, it is now common to perform synovectomies, especially of the knee, through the arthroscope. The arthroscope is a small, hollow rod inserted into the knee via a small puncture wound. In addition to permitting the intra-articular insertion of a fibre-optic system, the arthroscope allows access to surgical instruments, such that synovial tissue can be removed arthroscopically. Such procedures can be carried out under "spinal" anesthetic and the patient allowed home the same day. In this manner sufficient synovium can be obtained from patients who will receive this gene therapy.

10 The synovial cells (synoviocytes) contained within the excised tissue may be aseptically recovered by enzymic digestion of the connective tissue matrix. Generally, the synovium is cut into pieces of approximately 1 millimeter diameter and digested sequentially with trypsin (0.2% w/v in Grey's Balanced Salt Solution) for 30 minutes at 37° Centigrade, and collagenase (0.2% w/v in Grey's Balanced Salt Solution) for 2 hours at 37° Centigrade. Cells recovered from this digestion are seeded into plastic culture dishes at a concentration of 10^4 - 10^5 cells per square centimeter with Hank's F₁₂ medium supplemented with 10% fetal bovine serum and antibiotics. After 3-7 days, the culture medium is withdrawn. Non-adherent cells such as lymphocytes are removed by washing with Gey's Balanced Salt Solution and fresh medium added. The adherent cells can now be used as they are, allowed to grow to confluency or taken through one or more subcultures. Subculturing expands the cell number and removes non-dividing cells such as macrophages.

25 Following genetic manipulation of the cells thus recovered, they can be removed from the culture dish by trypsinizing, scraping or other means, and made into a standard suspension. Gey's Balanced Salt Solution or other isotonic salt solutions of suitable composition, or saline solution are suitable carriers. A suspension of cells can then be injected into the recipient mammalian joint. Intra-articular injections of this type are routine and easily carried out in the doctor's office. No surgery is necessary. Very large numbers of cells can be introduced in this way and repeat injections carried out as needed.

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Another embodiment of this invention is the gene produced by the hereinbefore described method of preparation. This gene carried by the retrovirus may be incorporated in a suitable pharmaceutical carrier, such as for example, buffered physiologic saline, for parenteral administration. This gene may be administered to a patient in a therapeutically effective dose.

5 More specifically, this gene may be incorporated in a suitable pharmaceutical carrier at a therapeutically effective dose and administered by intra-articular injection. Therefore, the preferred mode regarding the *ex vivo* method of delivery is the removal of the patients connective tissue (e.g., synovia), *in vitro* culture of this connective tissue, transduction of the DNA sequence of interest, followed by the above-mentioned manipulation prior to delivery to
10 the afflicted joint of the patient.

In another embodiment of this invention, this gene may be administered to patients as a prophylactic measure to prevent the development of arthritis in those patients determined to be highly susceptible of developing this disease. More specifically, this gene carried by the retrovirus may be incorporated in a suitable pharmaceutical carrier at a prophylactically effective
15 dose and administered by parenteral injection, including intra-articular injection.

EXAMPLE X

Fifty micrograms of a DNA plasmid vector molecule containing the interleukin-1 beta coding sequence ligated downstream of the CMV promoter was encapsulated within cationic liposomes, mixed with Geys biological buffer and injected intra-articularly into the knee
20 joints of a rabbit. Forty eight hours subsequent to injection one nanogram of interleukin-1 beta was recovered from the knee joint area. Therefore, injection of the DNA containing liposome solution within the region of the synovial tissue prompted fusion of the liposomes to the synovial cells, transfer of the DNA plasmid vector into synovial cells and subsequent expression of the IL-1 beta gene. Additionally, it is possible to inject non-encapsulated (i.e.,
25 naked) DNA into the joint area and monitor transfection of the DNA vector into the synovial cells as determined by subsequent expression of the IL-1 beta gene in synovial cells. Therefore, either method may be utilized as a plausible alternative to the *in vitro* manipulation of synovia also exemplified in the present invention.

EXAMPLE XI

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The *in vivo* biological activity of the MFG-IRAP construct was tested as the ability to suppress the effects of IL-1 β . Rabbit knees were injected with recombinant human IL-1 β , known to cause an increased concentration of leukocytes within the afflicted joint space. Introduction of MFG-IRAP/HIG-82 cells into rabbit knees strongly suppresses IL-1 β production of leukocytes to the afflicted joint space. In contrast, control HIG-82 cells do not suppress the leukocyte infiltration to the joint space challenged with IL-1 β (see Figure 10). Inhibition is greatest at the lowest doses of human recombinant IL-1 β (hrIL-1 β), as expected by the competitive mechanism through which IRAP antagonizes IL-1. Therefore, this rabbit model confirms that *in vivo*, intra-articular expression of IRAP is biologically active and can protect the joint from inflammation provoked by IL-1.

EXAMPLE XII

This example further evaluates *ex vivo* delivery into rabbit knee joints of the MFG-IRAP construct. As known, IRAP is an acidic glycoprotein of approximately 25 kDa that functions as a natural antagonist of the biological actions of interleukin-1 (IL-1) by binding to IL-1 receptors. Unlike IL-1, IRAP has no agonist activity, instead acting as a competitive inhibitor of the binding of IL-1.

This example shows that *in vivo* expression of IRAP by genetically modified synovial cells inhibits IL-1 β -induced leukocyte infiltration into the joint space, synovial thickening and hypercellularity, and loss of proteoglycans from articular cartilage.

As mentioned within this specification, the preferred mode of treating a patient through the *ex vivo* route will be by transplanting genetically modified autologous synovial cells by intra-articular injection. However, HIG-82 cells, easily maintained in culture, were used for these experiments to show that intra-articularly expressed IRAP is effective in inhibiting the physiological sequelae of intra-articularly injected IL-1.

MFG-IRAP/HIG-82 cells or control (uninfected HIG-82) cells, were transplanted into rabbit knees by intra-articular injection by the methods disclosed within this specification. Briefly, cultures of these cells were infected with MFG-IRAP. Media conditioned for 3 days by infected MFG-IRAP/HIG 82 cells were assayed for human IRAP by ELISA assay using a commercial kit (R&D Systems, Minneapolis, MN, USA) and found to contain approximately

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500 ng IRAP/ 10^6 cells. Western blotting confirmed the presence of human IRAP of size 22-25 kDa. HIG-IRAP cells were trypsinized, suspended in Gey's balanced salt solution and 1 ml of suspension, containing 10^7 cells, was injected intra-articularly into the left knee joints of New Zealand White rabbits (2.5 kg). The contralateral control knees received a similar injection of untransduced HIG-82 cells.

Three days following transplantation of the cells, knee joints were challenged by various doses of a single intra-articular injection of human recombinant IL-1 β dissolved in 0.5 ml Gey's solution. Control knees were injected with 0.5 ml of Gey's solution.

Eighteen hours after injection of hrIL-1 β , rabbits were killed and the knee joints evaluated histopathologically and for expression of IRAP. Each joint was first lavaged with 1 ml Gey's solution containing 10 mM EDTA. Cell counts in these washings were performed with a hemocytometer. An aliquot was removed for cytospinning and staining with 'DiffQuick' (Baxter Scientific Products) before examination under light microscopy. Washings were then centrifuged. Supernatants were removed for IRAP ELISA and for the determination of glycosaminoglycan (GAG) concentrations as an index of cartilage breakdown. GAG determinations were carried out with the dimethylmethylene blue assay (Farndale, et al., 1986, Biochim Biophys Acta 883: 173-177).

Synovia were dissected from the knee joints, fixed in 70% ethanol, dehydrated, embedded in paraffin, sectioned at 5 μ m and stained with hematoxylin and eosin.

An average of 2.5ng human IRAP per knee was measured in joint lavages 4 days following transplant of MFG-IRAP/HIG 82 cells. Contralateral, control knees receiving naive HIG-82 cells had no detectable human IRAP (Figure 11). To determine whether the observed level of IRAP expression was sufficient to inhibit the effects of IL-1 *in vivo*, increasing concentrations of IL-1 β (0-100 pg) were injected into both the control and IRAP knees. As is shown in Figure 12a, injection of hrIL-1 β into control knees provoked a marked leukocytosis which was strongly suppressed in the genetically modified knees. There was also a statistically significant reduction in the white blood cell count in knees containing MFG-IRAP/HIG 82 cells which had not been injected with IL-1. This may reflect the influence of IRAP upon the slight inflammatory effect of injecting cells into joints. The degree of suppression by IRAP decreased as the amount of injected hrIL-1 β increased, in

keeping with the competitive mode of inhibition existing between IRAP and IL-1. No dose-response for hrIL-1 β alone is evident in these particular experiments because this specific batch of IL-1 was especially effective in eliciting maximal response even at the lowest dose used.

5 Examination of cytopins (Figure 12b, 12c) revealed that most of the infiltrating leukocytes were neutrophils and monocytes. These preparations also serve to illustrate the efficiency with which leukocytosis was suppressed by the IRAP gene. Ten times the volume of lavage fluid is represented on the cytospin obtained from the IRAP-producing knees (Figure 12c) compared to the non-IRAP knees (Figure 12b).

10 To determine if intra-articularly expressed IRAP was able to block cartilage breakdown, the concentration of glycosaminoglycans (GAG) in joint lavages was determined. GAG analyses of the washings from the control and IRAP expressing knees (Figure 13) confirmed that transfer of the IRAP gene partially inhibited breakdown of the cartilaginous matrix in response to IL-1. Again, inhibition was strongest at the lowest concentrations of
15 IL-1 and was abolished at the highest dose of IL-1 (Figure 13).

 In response to 10 pg of injected hrIL-1 β , control synovia became hypertrophic (Figure 14a) and hypercellular (Figure 14c). The increased cellularity of the synovia appeared to involve both increased numbers of synoviocytes and infiltration by leukocytes. In knees where MFG-IRAP/HIG 82 cells were present, these changes were completely
20 suppressed and the synovia were nearly indistinguishable from control synovia (Figure 14b, 14d).

 The *ex vivo* transfer of the human IRAP gene to the synovial lining of rabbit knees clearly protects these joints from the pathophysiological sequelae of subsequent intra-articular challenge by hrIL-1 β .

25 Measurements of the amounts of IL-1 present in human, recombinant synovial fluids provide values in the range of 0-500 pg/ml (Westacott, et al., 1990, Ann Rheum Dis. 49: 676-681; Malvak, et al., 1993, Arthritis Rheum 36: 781-789). Thus the amounts of IRAP expressed intra-articularly during the present, short-term experiments should be sufficient to block the biological activities of IL-1 at the concentrations present in human arthritic joints.

30 EXAMPLE XIII

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This example shows that the level of intraarticular IRAP expressed subsequent to *ex vivo* transplantation of synoviocytes transduced with MFG-IRAP is sufficient to inhibit several pathophysiological changes associated with antigen-induced arthritis of the rabbit knee. Intraarticularly expressed IRAP has both a chondroprotective and anti-inflammatory effect during the acute phase of this disease. Data disclosed in Example XII support the contention that the invention as disclosed and claimed is a marked improvement for treating connective tissue disorders such as arthritis in comparison to delivery of proteins to the afflicted joint. Example XII shows that *ex vivo* transfer of MFG-IRAP to the rabbit knee as disclosed throughout this specification results in the intraarticular accumulation of nanogram quantities of glycosylated, biologically active IRAP. This present example shows that this same gene therapy based product inhibits joint pathologies in a rabbit model of human rheumatoid arthritis.

Young adult rabbits were subjected to a surgical, partial synovectomy of the left knee joint to provide autologous cells. These autologous cells were used to produce cultures of rabbit synovial fibroblasts (type B synoviocytes) from these biopsies as described in Example V and Example IX. Subconfluent cultures were then transduced by infection with MFG-IRAP. Expression of the transgene was confirmed by measuring the concentrations of human IRAP in the conditioned media; values typically range from 100-500ng IRAP/ 10^6 cells/3 days. Sister cultures of synoviocytes from the same animal were infected with a BAG virus encoding the lac Z and neo^r marker genes, and then selected for growth in the presence of G418 (1mg/ml) to serve as controls. Untransduced synoviocytes were also used as additional controls.

During the period that the cells were being grown and transduced, the donor rabbits were sensitized to ovalbumin by a series of two intradermal injections of 5mg ovalbumin emulsified in adjuvant, given two weeks apart. Two weeks after the second injection, an acute monarticular arthritis was initiated by the injection of 5mg ovalbumin dissolved in 1ml saline into the right knee joints. By this time the left, donor knees had regenerated their synovia, and were each injected with 1ml saline as controls.

One day after the onset of arthritis, 10^7 autologous cells, transduced with either the IRAP gene, or lac Z and neo genes, were injected into each arthritic knee, and each

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contralateral, non-arthritic knee. In other control experiments, knees were injected with untransduced, autologous cells. Groups of rabbits were killed 3 and 7 days later, corresponding to the middle and end of the acute phase of this arthropathy. Knees were lavaged with 1ml of saline, prior to the removal of synovial tissue and articular cartilage for analysis.

Intraarticular expression of the MFG-IRAP transgene was evaluated by ELISA measurements of human IRAP in the lavage fluids. IRAP concentrations in the control, non-arthritic knees is shown in Figure 15. IRAP concentrations in the arthritic knees were always several-fold higher than in normal knees at both time points (Figure 15). In both non-arthritic and arthritic knees transduced with MFG-IRAP, there was a slight decrease in IRAP expression with time. No human IRAP could be detected in sera obtained from normal or arthritic rabbits.

During the course of these experiments, the intraarticular concentration of rabbit IL-1 in arthritic knees was in the range of 100-200 pg/knee (Figure 16). No IL-1 α could be detected by RIA of the lavage fluids. Thus the concentration of IRAP within these knees exceeded the concentration of IL-1 by factors of approximately 10-50. Concentrations of IL-1 were lower in day 7 arthritic knees receiving the IRAP gene (Figure 16), suggesting that IRAP had inhibited an autocrine amplification loop.

Two major pathologies predominate in the rheumatoid joint: loss of articular cartilage and inflammation. The former occurs through a combination of reduced synthesis and enhanced degradation of the cartilaginous matrix. Whereas inflammation is manifest as a synovitis accompanied by influx of leukocytes into the joint space.

The onset of antigen-induced arthritis in this Example was accompanied by cartilage destruction, as reflected in the increased glycosaminoglycan (GAG) content of the lavage fluids (Figure 17a), and reduced synthesis of cartilage proteoglycans, as reflected by lower uptake of $^{35}\text{SO}_4^{2-}$ (Figure 17b). Knees expressing the MFG-IRAP transgene, but not control knees, were substantially protected from these changes. GAG release (Figure 17a) was inhibited 55% on day 4 and 32% on day 7. Suppression of GAG synthesis (Figure 17b) was inhibited by 68% on day 4 and 100% on day 7. The MFG-IRAP transgene also strongly reduced the influx of leukocytes into the joint space (Figure 18), an effect that was stronger

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at day 4 (65% inhibition) than at day 7 (38% inhibition); indeed, the difference at day 7 failed to reach statistical significance.

The MFG-IRAP construct is utilized to exemplify the presently claimed invention. In addition to this construct, the *ex vivo* based teachings of this specification have been utilized to transfer to synovial cells and express *in vivo* DNA sequences encoding human IL-1 α , human TNF- α soluble receptor, vIL-10, Lac Z and neo'.

EXAMPLE XIV

The methods disclosed throughout this specification were utilized to express MFG-human IL-1 soluble receptor type I and type II constructs (with neo') within *in vitro* cultured synoviocytes. These transfected synoviocytes produce 1-2 ng/10⁶ cells of IL-1 soluble receptor types I and II, following neo-selection. The additional methods disclosed throughout this specification may be utilized to procure *in vivo* expression data regarding these MFG-human IL-1 soluble receptor type I and type II constructs.

EXAMPLE XV

Rabbits were injected intraarticularly in one knee joint with a specific viral or non-viral vector disclosed in Table II. Contralateral knees were injected with a control, usually with the identical viral or non-viral vector with a different passenger gene. At intervals from 2 days to 2 weeks following intraarticular injection, rabbits were sacrificed and the knee joints harvested and stained with X-Gal to assay for LacZ expression. The results are depicted in Table II. The recombinant adenovirus vector comprising a CMV-LacZ fusion and the recombinant HSV vector comprising a CMV-LacZ fusion generated the highest expression level subsequent to intraarticular injection. The recombinant retroviral vector, MFG-LacZ, was not expressed *in vivo*, lending credence to the concept that retroviral vectors require actively dividing cells during the infection process and the concomitant low mitotic activity of synoviocytes in the joint lining.

However, an intra-articular injection of MFG-IRAP to synovial cells of an inflamed joint space supported retroviral transduction. Injection of MFG-IRAP into an inflamed rabbit knee lead to the intrarticular accumulation of 414.7 pg/knee at 24 hours post injection. The contralateral knee contained only 46.05 pg of human IRAP, close to background level.

TABLE II

VECTOR	PROMOTER	E X P R E S S I O N		DURATION (Days)
		In Vitro LAC Z cells (%)	In Vivo LEVEL	
Retrovirus (MFG)	LTR	20-30	0	0
HSV	CMV	1 (toxic)	+++	5-7
Adenovirus	CMV	100	+++	≥ 14
Liposome (DC-chol)	CMV	20-30	+	1-2
None (naked DNA)	CMV	0	±	1-2

Level of in vivo expression was evaluated subjectively on a scale of 0 - + + +, based upon the degree of staining with X-Gal.

LTR = viral long terminal repeat

CMV = cytomegalovirus

These data demonstrate that the gene therapy methods disclosed and claimed in the present invention can be used to modulate the disease process in an animal model of arthritis. In turn, these Examples enable the claimed gene therapy based treatment of connective tissue pathologies and systemic indices of inflammation within the afflicted joint(s). It will be appreciated by those skilled in the art that this invention provides a method of introducing into a connective tissue cell of a mammalian host *in vitro*, or in the alternative *in vivo*, at least one gene which codes for proteins with therapeutic properties. This method includes employing genes having DNA that is capable of maintenance and expression.

It will be appreciated by those skilled in the art that this invention provides a method of introducing at least one gene encoding a product into at least one cell of the connective tissue of a mammalian host for treating an arthritic condition of the mammalian host.

It will be understood by those skilled in the art that this invention provides a method to repair and regenerate the connective tissue of a mammalian host.

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It will be further understood that the present invention discloses *ex vivo* and *in vivo* techniques for delivery of a DNA sequence of interest to the connective tissue cells of the mammalian host. The *ex vivo* technique involves prior removal and culture of target autologous connective tissue cells, *in vitro* infection of the DNA sequence, DNA vector or other delivery vehicle of interest into the connective tissue cells, followed by transplantation to the modified connective tissue cells to the target joint of the mammalian host, so as to effect *in vivo* expression of the gene product of interest. The *in vivo* technique bypasses the requirement for *in vitro* culture of target connective tissues cells; instead relying on direct transplantation of the DNA sequence, DNA vector or other delivery vehicle to the target *in vivo* connective tissue cells, thus effecting expression of the gene product of interest.

It will be further understood that this invention provides a method to produce an animal model for the study of connective tissue pathology.

It will be appreciated by those persons skilled in the art that this invention provides a method of using and a method of preparing a gene encoding an extracellular interleukin-1 binding domain of an interleukin-1 receptor that is capable of binding to and neutralizing substantially all isoforms of interleukin-1, and thus substantially protect cartilage of a mammalian host from pathological degradation. In addition, it will be understood by those persons skilled in the art that the method of using the gene of this invention will reduce inflammation, protect soft tissues of the joint and suppress the loss of bone that occurs in patients suffering with arthritis.

It will be appreciated by those persons skilled in the art that the viral vectors employed in the hereinbefore described invention may be employed to transfect synovial cells *in vivo* or in culture, such as by direct intra-articular injection or transplantation of autologous synovial cells from the patient transduced with the retroviral vector carrying the truncated interleukin-1 receptor gene.

It will also be understood that a class of DNA sequences, as described throughout this specification, including but not limited to IRAP, may use the claimed methods to effect reduction of inflammation, protect soft tissues of the joint and suppress the loss of bone that occurs in patients suffering with arthritis.

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Whereas particular embodiments of this invention have been described above for purposes of illustration, it will be evident to those persons skilled in the art that numerous variations of the details of the present invention may be made without departing from the invention as defined in the appended claims.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT: University of Pittsburgh of the Commonwealth System of
Higher Education, 911 William Pitt Union
Pittsburgh, PA 15260

(ii) TITLE OF INVENTION: Gene Transfer For Treating a Connective Tissue of a
Mammalian Host

(iii) NUMBER OF SEQUENCES: 6

(iv) CORRESPONDENCE ADDRESS:

(A) ADDRESSEE: Eckert Seamans Cherin & Mellott
(B) STREET: 1700 Market Street Suite 3232
(C) CITY: Philadelphia
(D) STATE: PA
(E) COUNTRY: USA
(F) ZIP: 19103

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:
(B) FILING DATE:
(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Gould, Jr., Lewis F.
(B) REGISTRATION NUMBER: 25,057
(C) REFERENCE/DOCKET NUMBER: 109070-11A

(ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: (215) 575-6000
(B) TELEFAX: (215) 575-6015
(C) TELEX: 866172

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1770 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (iii) HYPOTHETICAL: NO
- (iv) ANTI-SENSE: NO
- (vii) IMMEDIATE SOURCE:
- (A) LIBRARY: Human T-cell cDNA Library
- (B) CLONE: Human Interleukin-1 Receptor
- (ix) FEATURE:
- (A) NAME/KEY: CDS
- (B) LOCATION: 55..1764

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

CCTCCTGAGA AGCTGGACCC CTTGGTAAAA GACAAGGCCT TCTCCAAGAA GAAT ATG	57
Met	
1	
AAA GTG TTA CTC AGA CTT ATT TGT TTC ATA GCT CTA CTG ATT TCT TCT	105
Lys Val Leu Leu Arg Leu Ile Cys Phe Ile Ala Leu Leu Ile Ser Ser	
5 10 15	
CTG GAG GCT GAT AAA TGC AAG GAA CGT GAA GAA AAA ATA ATT TTA GTG	153
Leu Glu Ala Asp Lys Cys Lys Glu Arg Glu Glu Lys Ile Ile Leu Val	
20 25 30	
TCA TCT GCA AAT GAA ATT GAT GTT CGT CCC TGT CCT CTT AAC CCA AAT	201
Ser Ser Ala Asn Glu Ile Asp Val Arg Pro Cys Pro Leu Asn Pro Asn	
35 40 45	
GAA CAC AAA GGC ACT ATA ACT TGG TAT AAA GAT GAC AGC AAG ACA CCT	249
Glu His Lys Gly Thr Ile Thr Trp Tyr Lys Asp Asp Ser Lys Thr Pro	
50 55 60 65	
GTA TCT ACA GAA CAA GCC TCC AGG ATT CAT CAA CAC AAA GAG AAA CTT	297
Val Ser Thr Glu Gln Ala Ser Arg Ile His Gln His Lys Glu Lys Leu	
70 75 80	
TGG TTT GTT CCT GCT AAG GTG GAG GAT TCA GGA CAT TAC TAT TGC GTG	345
Trp Phe Val Pro Ala Lys Val Glu Asp Ser Gly His Tyr Tyr Cys Val	
85 90 95	
GTA AGA AAT TCA TCT TAC TGC CTC AGA ATT AAA ATA AGT GCA AAA TTT	393
Val Arg Asn Ser Ser Tyr Cys Leu Arg Ile Lys Ile Ser Ala Lys Phe	
100 105 110	
GTG GAG AAT GAG CCT AAC TTA TGT TAT AAT GCA CAA GCC ATA TTT AAG	441
Val Glu Asn Glu Pro Asn Leu Cys Tyr Asn Ala Gln Ala Ile Phe Lys	
115 120 125	
CAG AAA CTA CCC GTT GCA GGA GAC GGA GGA CTT GTG TGC CCT TAT ATG	489
Gln Lys Leu Pro Val Ala Gly Asp Gly Gly Leu Val Cys Pro Tyr Met	
130 135 140 145	

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GAG	TTT	TTT	AAA	AAT	GAA	AAT	AAT	GAG	TTA	CCT	AAA	TTA	CAG	TGG	TAT	537
Glu	Phe	Phe	Lys	Asn	Glu	Asn	Asn	Glu	Leu	Pro	Lys	Leu	Gln	Trp	Tyr	
			150						155					160		
AAG	GAT	TGC	AAA	CCT	CTA	CTT	CTT	GAC	AAT	ATA	CAC	TTT	AGT	GGA	GTC	585
Lys	Asp	Cys	Lys	Pro	Leu	Leu	Leu	Asp	Asn	Ile	His	Phe	Ser	Gly	Val	
			165					170					175			
AAA	GAT	AGG	CTC	ATC	GTG	ATG	AAT	GTG	GCT	GAA	AAG	CAT	AGA	GGG	AAC	633
Lys	Asp	Arg	Leu	Ile	Val	Met	Asn	Val	Ala	Glu	Lys	His	Arg	Gly	Asn	
		180					185					190				
TAT	ACT	TGT	CAT	GCA	TCC	TAC	ACA	TAC	TTG	GGC	AAG	CAA	TAT	CCT	ATT	681
Tyr	Thr	Cys	His	Ala	Ser	Tyr	Thr	Tyr	Leu	Gly	Lys	Gln	Tyr	Pro	Ile	
	195					200					205					
ACC	CGG	GTA	ATA	GAA	TTT	ATT	ACT	CTA	GAG	GAA	AAC	AAA	CCC	ACA	AGG	729
Thr	Arg	Val	Ile	Glu	Phe	Ile	Thr	Leu	Glu	Glu	Asn	Lys	Pro	Thr	Arg	
210					215					220					225	
CCT	GTG	ATT	GTG	AGC	CCA	GCT	AAT	GAG	ACA	ATG	GAA	GTA	GAC	TTG	GGA	777
Pro	Val	Ile	Val	Ser	Pro	Ala	Asn	Glu	Thr	Met	Glu	Val	Asp	Leu	Gly	
				230					235					240		
TCC	CAG	ATA	CAA	TTG	ATC	TGT	AAT	GTC	ACC	GGC	CAG	TTG	AGT	GAC	ATT	825
Ser	Gln	Ile	Gln	Leu	Ile	Cys	Asn	Val	Thr	Gly	Gln	Leu	Ser	Asp	Ile	
			245					250					255			
GCT	TAC	TGG	AAG	TGG	AAT	GGG	TCA	GTA	ATT	GAT	GAA	GAT	GAC	CCA	GTG	873
Ala	Tyr	Trp	Lys	Trp	Asn	Gly	Ser	Val	Ile	Asp	Glu	Asp	Asp	Pro	Val	
		260				265						270				
CTA	GGG	GAA	GAC	TAT	TAC	AGT	GTG	GAA	AAT	CCT	GCA	AAC	AAA	AGA	AGG	921
Leu	Gly	Glu	Asp	Tyr	Tyr	Ser	Val	Glu	Asn	Pro	Ala	Asn	Lys	Arg	Arg	
	275					280					285					
AGT	ACC	CTC	ATC	ACA	GTG	CTT	AAT	ATA	TCG	GAA	ATT	GAA	AGT	AGA	TTT	969
Ser	Thr	Leu	Ile	Thr	Val	Leu	Asn	Ile	Ser	Glu	Ile	Glu	Ser	Arg	Phe	
290					295					300					305	
TAT	AAA	CAT	CCA	TTT	ACC	TGT	TTT	GCC	AAG	AAT	ACA	CAT	GGT	ATA	GAT	1017
Tyr	Lys	His	Pro	Phe	Thr	Cys	Phe	Ala	Lys	Asn	Thr	His	Gly	Ile	Asp	
			310						315					320		
GCA	GCA	TAT	ATC	CAG	TTA	ATA	TAT	CCA	GTC	ACT	AAT	TTC	CAG	AAG	CAC	1065
Ala	Ala	Tyr	Ile	Gln	Leu	Ile	Tyr	Pro	Val	Thr	Asn	Phe	Gln	Lys	His	
			325					330					335			
ATG	ATT	GGT	ATA	TGT	GTC	ACG	TTG	ACA	GTC	ATA	ATT	GTG	TGT	TCT	GTT	1113
Met	Ile	Gly	Ile	Cys	Val	Thr	Leu	Thr	Val	Ile	Ile	Val	Cys	Ser	Val	
		340					345					350				
TTC	ATC	TAT	AAA	ATC	TTC	AAG	ATT	GAC	ATT	GTG	CTT	TGG	TAC	AGG	GAT	1161
Phe	Ile	Tyr	Lys	Ile	Phe	Lys	Ile	Asp	Ile	Val	Leu	Trp	Tyr	Arg	Asp	
		355				360					365					
TCC	TGC	TAT	GAT	TTT	CTC	CCA	ATA	AAA	GCT	TCA	GAT	GGA	AAG	ACC	TAT	1209
Ser	Cys	Tyr	Asp	Phe	Leu	Pro	Ile	Lys	Ala	Ser	Asp	Gly	Lys	Thr	Tyr	
370					375					380					385	
GAC	GCA	TAT	ATA	CTG	TAT	CCA	AAG	ACT	GTT	GGG	GAA	GGG	TCT	ACC	TCT	1257
Asp	Ala	Tyr	Ile	Leu	Tyr	Pro	Lys	Thr	Val	Gly	Glu	Gly	Ser	Thr	Ser	
			390						395					400		
GAC	TGT	GAT	ATT	TTT	GTG	TTT	AAA	GTC	TTG	CCT	GAG	GTC	TTG	GAA	AAA	1305
Asp	Cys	Asp	Ile	Phe	Val	Phe	Lys	Val	Leu	Pro	Glu	Val	Leu	Glu	Lys	
			405					410					415			

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CAG TGT GGA TAT AAG CTG TTC ATT TAT GGA AGG GAT GAC TAC GTT GGG Gln Cys Gly Tyr Lys Leu Phe Ile Tyr Gly Arg Asp Asp Tyr Val Gly 420 425 430	1353
GAA GAC ATT GTT GAG GTC ATT AAT GAA AAC GTA AAG AAA AGC AGA AGA Glu Asp Ile Val Glu Val Ile Asn Glu Asn Val Lys Lys Ser Arg Arg 435 440 445	1401
CTG ATT ATC ATT TTA GTC AGA GAA ACA TCA GGC TTC AGC TGG CTG GGT Leu Ile Ile Ile Leu Val Arg Glu Thr Ser Gly Phe Ser Trp Leu Gly 450 455 460 465	1449
GGT TCA TCT GAA GAG CAA ATA GCC ATG TAT AAT GCT CTT GTT CAG GAT Gly Ser Ser Glu Glu Gln Ile Ala Met Tyr Asn Ala Leu Val Gln Asp 470 475 480	1497
GGA ATT AAA GTT GTC CTG CTT GAG CTG GAG AAA ATC CAA GAC TAT GAG Gly Ile Lys Val Val Leu Leu Glu Leu Glu Lys Ile Gln Asp Tyr Glu 485 490 495	1545
AAA ATG CCA GAA TCG ATT AAA TTC ATT AAG CAG AAA CAT GGG GCT ATC Lys Met Pro Glu Ser Ile Lys Phe Ile Lys Gln Lys His Gly Ala Ile 500 505 510	1593
CGC TGG TCA GGG GAC TTT ACA CAG GGA CCA CAG TCT GCA AAG ACA AGG Arg Trp Ser Gly Asp Phe Thr Gln Gly Pro Gln Ser Ala Lys Thr Arg 515 520 525	1641
TTC TGG AAG AAT GTC AGG TAC CAC ATG CCA GTC CAG CGA CGG TCA CCT Phe Trp Lys Asn Val Arg Tyr His Met Pro Val Gln Arg Arg Ser Pro 530 535 540 545	1689
TCA TCT AAA CAC CAG TTA CTG TCA CCA GCC ACT AAG GAG AAA CTG CAA Ser Ser Lys His Gln Leu Leu Ser Pro Ala Thr Lys Glu Lys Leu Gln 550 555 560	1737
AGA GAG GCT CAC GTG CCT CTC GGG TAGCATGGA Arg Glu Ala His Val Pro Leu Gly 565 570	1770

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 569 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met	Lys	Val	Leu	Leu	Arg	Leu	Ile	Cys	Phe	Ile	Ala	Leu	Leu	Ile	Ser	1	5	10	15
Ser	Leu	Glu	Ala	Asp	Lys	Cys	Lys	Glu	Arg	Glu	Glu	Lys	Ile	Ile	Leu	20	25	30	
Val	Ser	Ser	Ala	Asn	Glu	Ile	Asp	Val	Arg	Pro	Cys	Pro	Leu	Asn	Pro	35	40	45	

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Asn Glu His Lys Gly Thr Ile Thr Trp Tyr Lys Asp Asp Ser Lys Thr
 50 55 60
 Pro Val Ser Thr Glu Gln Ala Ser Arg Ile His Gln His Lys Glu Lys
 65 70 75 80
 Leu Trp Phe Val Pro Ala Lys Val Glu Asp Ser Gly His Tyr Tyr Cys
 85 90 95
 Val Val Arg Asn Ser Ser Tyr Cys Leu Arg Ile Lys Ile Ser Ala Lys
 100 105 110
 Phe Val Glu Asn Glu Pro Asn Leu Cys Tyr Asn Ala Gln Ala Ile Phe
 115 120 125
 Lys Gln Lys Leu Pro Val Ala Gly Asp Gly Gly Leu Val Cys Pro Tyr
 130 135 140
 Met Glu Phe Phe Lys Asn Glu Asn Asn Glu Leu Pro Lys Leu Gln Trp
 145 150 155 160
 Tyr Lys Asp Cys Lys Pro Leu Leu Leu Asp Asn Ile His Phe Ser Gly
 165 170 175
 Val Lys Asp Arg Leu Ile Val Met Asn Val Ala Glu Lys His Arg Gly
 180 185 190
 Asn Tyr Thr Cys His Ala Ser Tyr Thr Tyr Leu Gly Lys Gln Tyr Pro
 195 200 205
 Ile Thr Arg Val Ile Glu Phe Ile Thr Leu Glu Glu Asn Lys Pro Thr
 210 215 220
 Arg Pro Val Ile Val Ser Pro Ala Asn Glu Thr Met Glu Val Asp Leu
 225 230 235 240
 Gly Ser Gln Ile Gln Leu Ile Cys Asn Val Thr Gly Gln Leu Ser Asp
 245 250 255
 Ile Ala Tyr Trp Lys Trp Asn Gly Ser Val Ile Asp Glu Asp Asp Pro
 260 265 270
 Val Leu Gly Glu Asp Tyr Tyr Ser Val Glu Asn Pro Ala Asn Lys Arg
 275 280 285
 Arg Ser Thr Leu Ile Thr Val Leu Asn Ile Ser Glu Ile Glu Ser Arg
 290 295 300
 Phe Tyr Lys His Pro Phe Thr Cys Phe Ala Lys Asn Thr His Gly Ile
 305 310 315 320
 Asp Ala Ala Tyr Ile Gln Leu Ile Tyr Pro Val Thr Asn Phe Gln Lys
 325 330 335
 His Met Ile Gly Ile Cys Val Thr Leu Thr Val Ile Ile Val Cys Ser
 340 345 350
 Val Phe Ile Tyr Lys Ile Phe Lys Ile Asp Ile Val Leu Trp Tyr Arg
 355 360 365
 Asp Ser Cys Tyr Asp Phe Leu Pro Ile Lys Ala Ser Asp Gly Lys Thr
 370 375 380
 Tyr Asp Ala Tyr Ile Leu Tyr Pro Lys Thr Val Gly Glu Gly Ser Thr
 385 390 395 400

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Ser Asp Cys Asp Ile Phe Val Phe Lys Val Leu Pr Glu Val Leu Glu
 405 410 415
 Lys Gln Cys Gly Tyr Lys Leu Phe Ile Tyr Gly Arg Asp Asp Tyr Val
 420 425 430
 Gly Glu Asp Ile Val Glu Val Ile Asn Glu Asn Val Lys Lys Ser Arg
 435 440 445
 Arg Leu Ile Ile Ile Leu Val Arg Glu Thr Ser Gly Phe Ser Trp Leu
 450 455 460
 Gly Gly Ser Ser Glu Glu Gln Ile Ala Met Tyr Asn Ala Leu Val Gln
 465 470 475 480
 Asp Gly Ile Lys Val Val Leu Leu Glu Leu Glu Lys Ile Gln Asp Tyr
 485 490 495
 Glu Lys Met Pro Glu Ser Ile Lys Phe Ile Lys Gln Lys His Gly Ala
 500 505 510
 Ile Arg Trp Ser Gly Asp Phe Thr Gln Gly Pro Gln Ser Ala Lys Thr
 515 520 525
 Arg Phe Trp Lys Asn Val Arg Tyr His Met Pro Val Gln Arg Arg Ser
 530 535 540
 Pro Ser Ser Lys His Gln Leu Leu Ser Pro Ala Thr Lys Glu Lys Leu
 545 550 555 560
 Gln Arg Glu Ala His Val Pro Leu Gly
 565

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1782 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(vii) IMMEDIATE SOURCE:

- (A) LIBRARY: Mouse T-cell cDNA Library
- (B) CLONE: Mouse Interleukin-1 Receptor

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 46..1776

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GGATGTCATC AGAGTTCCCA GTGCCCCGAA CCGTGAACAA CACAA ATG GAG AAT	54
Met Glu Asn	
1	
ATG AAA GTG CTA CTG GGG CTC ATT TGT CTC ATG GTG CCT CTG CTG TCG	102
Met Lys Val Leu Leu Gly Leu Ile Cys Leu Met Val Pro Leu Leu Ser	
5 10 15	
CTG GAG ATT GAC GTA TGT ACA GAA TAT CCA AAT CAG ATC GTT TTG TTT	150
Leu Glu Ile Asp Val Cys Thr Glu Tyr Pro Asn Gln Ile Val Leu Phe	
20 25 30 35	
TTA TCT GTA AAT GAA ATT GAT ATT CGC AAG TGT CCT CTT ACT CCA AAT	198
Leu Ser Val Asn Glu Ile Asp Ile Arg Lys Cys Pro Leu Thr Pro Asn	
40 45 50	
AAA ATG CAC GGC GAC ACC ATA ATT TGG TAC AAG AAT GAC AGC AAG ACC	246
Lys Met His Gly Asp Thr Ile Ile Trp Tyr Lys Asn Asp Ser Lys Thr	
55 60 65	
CCC ATA TCA GCG GAC CGG GAC TCC AGG ATT CAT CAG CAG AAT GAA CAT	294
Pro Ile Ser Ala Asp Arg Asp Ser Arg Ile His Gln Gln Asn Glu His	
70 75 80	
CTT TGG TTT GTA CCT GCC AAG GTG GAG GAC TCA GGA TAT TAC TAT TGT	342
Leu Trp Phe Val Pro Ala Lys Val Glu Asp Ser Gly Tyr Tyr Tyr Cys	
85 90 95	
ATA GTA AGA AAC TCA ACT TAC TGC CTC AAA ACT AAA GTA ACC GTA ACT	390
Ile Val Arg Asn Ser Thr Tyr Cys Leu Lys Thr Lys Val Thr Val Thr	
100 105 110 115	
GTG TTA GAG AAT GAC CCT GGC TTG TGT TAC AGC ACA CAG GCC ACC TTC	438
Val Leu Glu Asn Asp Pro Gly Leu Cys Tyr Ser Thr Gln Ala Thr Phe	
120 125 130	
CCA CAG CGG CTC CAC ATT GCC GGG GAT GGA AGT CTT GTG TGC CCT TAT	486
Pro Gln Arg Leu His Ile Ala Gly Asp Gly Ser Leu Val Cys Pro Tyr	
135 140 145	
GTG AGT TAT TTT AAA GAT GAA AAT AAT GAG TTA CCC GAG GTC CAG TGG	534
Val Ser Tyr Phe Lys Asp Glu Asn Asn Glu Leu Pro Glu Val Gln Trp	
150 155 160	
TAT AAG AAC TGT AAA CCT CTG CTT CTT GAC AAC GTG AGC TTC TTC GGA	582
Tyr Lys Asn Cys Lys Pro Leu Leu Leu Asp Asn Val Ser Phe Phe Gly	
165 170 175	
GTA AAA GAT AAA CTG TTG GTG AGG AAT GTG GCT GAA GAG CAC AGA GGG	630
Val Lys Asp Lys Leu Leu Val Arg Asn Val Ala Glu Glu His Arg Gly	
180 185 190 195	
GAC TAT ATA TGC CGT ATG TCC TAT ACG TTC CGG GGG AAG CAA TAT CCG	678
Asp Tyr Ile Cys Arg Met Ser Tyr Thr Phe Arg Gly Lys Gln Tyr Pro	
200 205 210	
GTC ACA CGA GTA ATA CAA TTT ATC ACA ATA GAT GAA AAC AAG AGG GAC	726
Val Thr Arg Val Ile Gln Phe Ile Thr Ile Asp Glu Asn Lys Arg Asp	
215 220 225	
AGA CCT GTT ATC CTG AGC CCT CGG AAT GAG ACG ATC GAA GCT GAC CCA	774
Arg Pro Val Ile Leu Ser Pro Arg Asn Glu Thr Ile Glu Ala Asp Pro	
230 235 240	

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GGA	TCA	ATG	ATA	CAA	CTG	ATC	TGC	AAC	GTC	ACG	GGC	CAG	TTC	TCA	GAC	822
Gly	Ser	M t	Ile	Gln	Leu	Ile	Cys	Asn	Val	Thr	Gly	Gln	Phe	Ser	Asp	
245						250					255					
CTT	GTC	TAC	TGG	AAG	TGG	AAT	GGA	TCA	GAA	ATT	GAA	TGG	AAT	GAT	CCA	870
Leu	Val	Tyr	Trp	Lys	Trp	Asn	Gly	Ser	Glu	Ile	Glu	Trp	Asn	Asp	Pro	
260				265						270					275	
TTT	CTA	GCT	GAA	GAC	TAT	CAA	TTT	GTG	GAA	CAT	CCT	TCA	ACC	AAA	AGA	918
Phe	Leu	Ala	Glu	Asp	Tyr	Gln	Phe	Val	Glu	His	Pro	Ser	Thr	Lys	Arg	
			280						285					290		
AAA	TAC	ACA	CTC	ATT	ACA	ACA	CTT	AAC	ATT	TCA	GAA	GTT	AAA	AGC	CAG	966
Lys	Tyr	Thr	Leu	Ile	Thr	Thr	Leu	Asn	Ile	Ser	Glu	Val	Lys	Ser	Gln	
			295					300					305			
TTT	TAT	CGC	TAT	CCG	TTT	ATC	TGT	GTT	GTT	AAG	AAC	ACA	AAT	ATT	TTT	1014
Phe	Tyr	Arg	Tyr	Pro	Phe	Ile	Cys	Val	Val	Lys	Asn	Thr	Asn	Ile	Phe	
		310					315					320				
GAG	TCG	GCG	CAT	GTG	CAG	TTA	ATA	TAC	CCA	GTC	CCT	GAC	TTC	AAG	AAT	1062
Glu	Ser	Ala	His	Val	Gln	Leu	Ile	Tyr	Pro	Val	Pro	Asp	Phe	Lys	Asn	
	325					330					335					
TAC	CTC	ATC	GGG	GGC	TTT	ATC	ATC	CTC	ACG	GCT	ACA	ATT	GTA	TGC	TGT	1110
Tyr	Leu	Ile	Gly	Gly	Phe	Ile	Ile	Leu	Thr	Ala	Thr	Ile	Val	Cys	Cys	
340					345				350						355	
GTG	TGC	ATC	TAT	AAA	GTC	TTC	AAG	GTT	GAC	ATA	GTG	CTT	TGG	TAC	AGG	1158
Val	Cys	Ile	Tyr	Lys	Val	Phe	Lys	Val	Asp	Ile	Val	Leu	Trp	Tyr	Arg	
				360					365					370		
GAC	TCC	TGC	TCT	GGT	TTT	CTT	CCT	TCA	AAA	GCT	TCA	GAT	GGA	AAG	ACA	1206
Asp	Ser	Cys	Ser	Gly	Phe	Leu	Pro	Ser	Lys	Ala	Ser	Asp	Gly	Lys	Thr	
			375					380					385			
TAC	GAT	GCA	TAT	ATT	CTT	TAT	CCC	AAG	ACC	CTG	GGA	GAG	GGG	TCC	TTC	1254
Tyr	Asp	Ala	Tyr	Ile	Leu	Tyr	Pro	Lys	Thr	Leu	Gly	Glu	Gly	Ser	Phe	
		390					395					400				
TCA	GAC	TTA	GAT	ACT	TTT	GTT	TTT	AAA	CTG	TTG	CCT	GAG	GTC	TTG	GAG	1302
Ser	Asp	Leu	Asp	Thr	Phe	Val	Phe	Lys	Leu	Leu	Pro	Glu	Val	Leu	Glu	
	405					410					415					
GGA	CAG	TTT	GGA	TAC	AAG	CTG	TTC	ATT	TAT	GGA	AGG	GAT	GAC	TAT	GTT	1350
Gly	Gln	Phe	Gly	Tyr	Lys	Leu	Phe	Ile	Tyr	Gly	Arg	Asp	Asp	Tyr	Val	
420					425					430					435	
GGA	GAA	GAT	ACC	ATC	GAG	GTT	ACT	AAT	GAA	AAT	GTA	AAG	AAA	AGC	AGG	1398
Gly	Glu	Asp	Thr	Ile	Glu	Val	Thr	Asn	Glu	Asn	Val	Lys	Lys	Ser	Arg	
				440					445					450		
AGG	CTG	ATT	ATC	ATT	CTA	GTG	AGA	GAT	ATG	GGA	GGC	TTC	AGC	TGG	CTG	1446
Arg	Leu	Ile	Ile	Ile	Leu	Val	Arg	Asp	Met	Gly	Gly	Phe	Ser	Trp	Leu	
			455					460					465			
GGC	CAG	TCA	TCT	GAA	GAG	CAA	ATA	GCC	ATA	TAC	AAT	GCT	CTC	ATC	CAG	1494
Gly	Gln	Ser	Ser	Glu	Glu	Gln	Ile	Ala	Ile	Tyr	Asn	Ala	Leu	Ile	Gln	
		470					475					480				
GAA	GGA	ATT	AAA	ATC	GTC	CTG	CTT	GAG	TTG	GAG	AAA	ATC	CAA	GAC	TAT	1542
Glu	Gly	Ile	Lys	Ile	Val	Leu	Leu	Glu	Leu	Glu	Lys	Ile	Gln	Asp	Tyr	
		485				490					495					
GAG	AAA	ATG	CCA	GAT	TCT	ATT	CAG	TTC	ATT	AAG	CAG	AAA	CAC	GGA	GTC	1590
Glu	Lys	Met	Pro	Asp	Ser	Ile	Gln	Phe	Ile	Lys	Gln	Lys	His	Gly	Val	
500					505					510					515	

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ATT TGC TGG TCA GGA GAC TTT CAA GAA AGA CCA CAG TCT GCA AAG ACC	1638
Ile Cys Trp Ser Gly Asp Phe Gln Glu Arg Pro Gln Ser Ala Lys Thr	
520 525 530	
AGG TTC TGG AAA AAC TTA AGA TAC CAG ATG CCA GCC CAA CGG AGA TCA	1686
Arg Phe Trp Lys Asn Leu Arg Tyr Gln Met Pro Ala Gln Arg Arg Ser	
535 540 545	
CCA TTG TCT AAA CAC CGC TTA CTA ACC CTG GAT CCT GTG CGG GAC ACT	1734
Pro Leu Ser Lys His Arg Leu Leu Thr Leu Asp Pro Val Arg Asp Thr	
550 555 560	
AAG GAG AAA CTG CCG GCA GCA ACA CAC TTA CCA CTC GGC TAGCATGGC	1782
Lys Glu Lys Leu Pro Ala Ala Thr His Leu Pro Leu Gly	
565 570 575	

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 576 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Met	Glu	Asn	Met	Lys	Val	Leu	Leu	Gly	Leu	Ile	Cys	Leu	Met	Val	Pro
1				5					10					15	
Leu	Leu	Ser	Leu	Glu	Ile	Asp	Val	Cys	Thr	Glu	Tyr	Pro	Asn	Gln	Ile
			20					25					30		
Val	Leu	Phe	Leu	Ser	Val	Asn	Glu	Ile	Asp	Ile	Arg	Lys	Cys	Pro	Leu
		35				40						45			
Thr	Pro	Asn	Lys	Met	His	Gly	Asp	Thr	Ile	Ile	Trp	Tyr	Lys	Asn	Asp
		50				55					60				
Ser	Lys	Thr	Pro	Ile	Ser	Ala	Asp	Arg	Asp	Ser	Arg	Ile	His	Gln	Gln
	65				70					75				80	
Asn	Glu	His	Leu	Trp	Phe	Val	Pro	Ala	Lys	Val	Glu	Asp	Ser	Gly	Tyr
			85						90					95	
Tyr	Tyr	Cys	Ile	Val	Arg	Asn	Ser	Thr	Tyr	Cys	Leu	Lys	Thr	Lys	Val
		100						105					110		
Thr	Val	Thr	Val	Leu	Glu	Asn	Asp	Pro	Gly	Leu	Cys	Tyr	Ser	Thr	Gln
		115					120					125			
Ala	Thr	Phe	Pro	Gln	Arg	Leu	His	Ile	Ala	Gly	Asp	Gly	Ser	Leu	Val
		130				135					140				
Cys	Pro	Tyr	Val	Ser	Tyr	Phe	Lys	Asp	Glu	Asn	Asn	Glu	Leu	Pro	Glu
	145				150					155				160	
Val	Gln	Trp	Tyr	Lys	Asn	Cys	Lys	Pro	Leu	Leu	Leu	Asp	Asn	Val	Ser
			165					170						175	
Phe	Phe	Gly	Val	Lys	Asp	Lys	Leu	Leu	Val	Arg	Asn	Val	Ala	Glu	Glu

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180	185	190
His Arg Gly Asp Tyr Ile Cys Arg Met Ser Tyr Thr Phe Arg Gly Lys 195 200 205		
Gln Tyr Pro Val Thr Arg Val Ile Gln Phe Ile Thr Ile Asp Glu Asn 210 215 220		
Lys Arg Asp Arg Pro Val Ile Leu Ser Pro Arg Asn Glu Thr Ile Glu 225 230 235 240		
Ala Asp Pro Gly Ser Met Ile Gln Leu Ile Cys Asn Val Thr Gly Gln 245 250 255		
Phe Ser Asp Leu Val Tyr Trp Lys Trp Asn Gly Ser Glu Ile Glu Trp 260 265 270		
Asn Asp Pro Phe Leu Ala Glu Asp Tyr Gln Phe Val Glu His Pro Ser 275 280 285		
Thr Lys Arg Lys Tyr Thr Leu Ile Thr Thr Leu Asn Ile Ser Glu Val 290 295 300		
Lys Ser Gln Phe Tyr Arg Tyr Pro Phe Ile Cys Val Val Lys Asn Thr 305 310 315 320		
Asn Ile Phe Glu Ser Ala His Val Gln Leu Ile Tyr Pro Val Pro Asp 325 330 335		
Phe Lys Asn Tyr Leu Ile Gly Gly Phe Ile Ile Leu Thr Ala Thr Ile 340 345 350		
Val Cys Cys Val Cys Ile Tyr Lys Val Phe Lys Val Asp Ile Val Leu 355 360 365		
Trp Tyr Arg Asp Ser Cys Ser Gly Phe Leu Pro Ser Lys Ala Ser Asp 370 375 380		
Gly Lys Thr Tyr Asp Ala Tyr Ile Leu Tyr Pro Lys Thr Leu Gly Glu 385 390 395 400		
Gly Ser Phe Ser Asp Leu Asp Thr Phe Val Phe Lys Leu Leu Pro Glu 405 410 415		
Val Leu Glu Gly Gln Phe Gly Tyr Lys Leu Phe Ile Tyr Gly Arg Asp 420 425 430		
Asp Tyr Val Gly Glu Asp Thr Ile Glu Val Thr Asn Glu Asn Val Lys 435 440 445		
Lys Ser Arg Arg Leu Ile Ile Ile Leu Val Arg Asp Met Gly Gly Phe 450 455 460		
Ser Trp Leu Gly Gln Ser Ser Glu Glu Gln Ile Ala Ile Tyr Asn Ala 465 470 475 480		
Leu Ile Gln Glu Gly Ile Lys Ile Val Leu Leu Glu Leu Glu Lys Ile 485 490 495		
Gln Asp Tyr Glu Lys Met Pro Asp Ser Ile Gln Phe Ile Lys Gln Lys 500 505 510		
His Gly Val Ile Cys Trp Ser Gly Asp Phe Gln Glu Arg Pro Gln Ser 515 520 525		
Ala Lys Thr Arg Phe Trp Lys Asn L u Arg Tyr Gln Met Pr Ala Gln 530 535 540		

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Arg	Arg	Ser	Pr	Leu	Ser	Lys	His	Arg	Leu	Leu	Thr	Leu	Asp	Pro	Val
545					550					555					560
Arg	Asp	Thr	Lys	Glu	Lys	Leu	Pro	Ala	Ala	Thr	His	Leu	Pro	Leu	Gly
				565					570					575	

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iii) HYPOTHETICAL: NO
- (iv) ANTI-SENSE: NO
- (vii) IMMEDIATE SOURCE:
 - (B) CLONE: Primer Oligonucleotide to 5'Leader Sequence of IL-1 Receptor
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

GCGGATCCCC TCCTGAGAAG CT

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(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iii) HYPOTHETICAL: NO
- (iv) ANTI-SENSE: NO
- (vii) IMMEDIATE SOURCE:
 - (B) CLONE: Primer Oligonucleotide Upstream of Transmembrane Portion of
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

GCGGATCCCA TGTGCTACTG G

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WHAT IS CLAIMED IS:

1. A method of treating a connective tissue disorder which comprises:
 - a) generating a recombinant viral vector which contains a DNA sequence encoding a protein or biologically active fragment thereof;
 - b) infecting a population of *in vitro* cultured connective tissue cells with said recombinant viral vector, resulting in a population of transfected connective cells; and,
 - c) transplanting said transfected connective cells to a joint space of a mammalian host, such that subsequent expression of said protein within said joint space substantially reduces at least one deleterious joint pathology or indicia of inflammation normally associated with a connective tissue disorder.
2. The method of claim 1 wherein said connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament.
3. The method of claim 2 wherein said connective tissue cells are synovial cells.
4. The method of claim 3 wherein said recombinant viral vector is a retroviral vector.
5. The method of claim 4 wherein transplantation of transduced synovial cells is by intraarticular injection.
6. The method of claim 5 wherein said synovial cells are autologous synovial cells.
7. The method of claim 4 wherein said recombinant retroviral vector comprises a DNA sequence encoding IRAP or a biologically active fragment thereof.
8. The method of claim 5 wherein said recombinant retroviral vector comprises a DNA sequence encoding IRAP or a biologically active fragment thereof.
9. The method of claim 6 wherein said recombinant retroviral vector comprises a DNA sequence encoding IRAP or a biologically active fragment thereof.
10. The method of claim 4 wherein said recombinant retroviral vector is MFG-IRAP.
11. The method of claim 5 wherein said recombinant retroviral vector is MFG-IRAP.

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12. The method of claim 6 wherein said recombinant retroviral vector is MFG-IRAP.
13. The method of claim 4 wherein said recombinant retroviral vector comprises a DNA sequence encoding the extracellular domain of the interleukin-1 receptor protein or a biologically active fragment thereof.
14. The method of claim 13 wherein said retroviral vector is MFG.
15. The method of claim 4 wherein said recombinant retroviral vector comprises a DNA sequence encoding the TNF- α soluble receptor protein or a biologically active fragment thereof.
16. The method of claim 15 wherein said retroviral vector is MFG.
17. The method of claim 4 wherein said recombinant retroviral vector comprises a DNA sequence encoding the IL4 protein or a biologically active fragment thereof.
18. The method of claim 17 wherein said retroviral vector is MFG.
19. The method of claim 4 wherein said recombinant retroviral vector comprises a DNA sequence encoding the IL10 protein or a biologically active fragment thereof.
20. The method of claim 19 wherein said retroviral vector is MFG.
21. The method of claim 1 wherein said population of transfected connective cells are stored prior to transplantation.
22. The method of claim 21 wherein said population of transfected connective cells are stored in 10% DMSO under liquid nitrogen prior to transplantation.
23. The method of claim 2 wherein said connective tissue is employed to regenerate damaged connective tissue within the joint space.
24. A method of treating a connective tissue disorder which comprises:
 - a) generating a recombinant DNA plasmid vector which contains a DNA sequence encoding a protein or biologically active fragment thereof;
 - b) transforming a population of *in vitro* cultured connective tissue cells with said recombinant DNA plasmid vector, resulting in a population of transformed connective cells; and,
 - c) transplanting said transformed connective cells to a joint space of a mammalian host, such that subsequent expression of said protein within said joint space

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substantially reduces at least one deleterious joint pathology or indicia of inflammation normally associated with a connective tissue disorder.

25. The method of claim 24 wherein transforming means is selected from the group consisting of liposome encapsulation, calcium phosphate, electroporation and DEAE-dextran.

26. The method of claim 24 wherein said connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament.

27. A method to produce an animal model for the study of joint pathologies which comprises:

a) generating a recombinant viral vector which contains a DNA sequence encoding a protein or biologically active fragment thereof;

b) infecting a population of *in vitro* cultured connective tissue cells with said recombinant viral vector, resulting in a population of transfected connective cells; and,

c) transplanting said transfected connective cells to a joint space of a mammalian host.

28. The method of claim 27 wherein said connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament.

29. The method of claim 28 wherein said connective tissue cells are synovial cells.

30. The method of claim 29 wherein said recombinant viral vector is a retroviral vector.

31. The method of claim 30 wherein transplantation of transduced synovial cells is by intraarticular injection.

32. The method of claim 31 wherein said synovial cells are autologous synovial cells.

33. The method of claim 32 wherein said recombinant retroviral vector comprises a DNA sequence encoding human IL-1 α or a biologically active fragment thereof.

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34. The method of claim 32 wherein said recombinant retroviral vector comprises a DNA sequence encoding human IL-1 β or a biologically active fragment thereof.

35. The method of claim 32 wherein said recombinant retroviral vector comprises a DNA sequence encoding human tumor necrosis factor- α or a biologically active fragment thereof.

36. A method to produce an animal model for the study of joint pathologies which comprises:

a) generating a recombinant DNA plasmid vector which contains a DNA sequence encoding a protein or biologically active fragment thereof;

b) transforming a population of *in vitro* cultured connective tissue cells with said recombinant DNA plasmid vector, resulting in a population of transformed connective cells; and,

c) transplanting said transformed connective cells to a joint space of a mammalian host.

37. The method of claim 36 wherein said connective tissue cells are selected from the group of connective tissue consisting of a synovium, a cartilage, a tendon and a ligament.

38. The method of claim 37 wherein said connective tissue cells are synovial cells.

39. The method of claim 38 wherein said recombinant plasmid DNA vector comprises a DNA sequence encoding human IL-1 α or a biologically active fragment thereof.

40. The method of claim 38 wherein said recombinant plasmid DNA vector comprises a DNA sequence encoding human IL-1 β or a biologically active fragment thereof.

41. The method of claim 38 wherein said recombinant plasmid DNA vector comprises a DNA sequence encoding human tumor necrosis factor- α or a biologically active fragment thereof.

42. The method of claim 40 wherein said recombinant plasmid DNA vector is CMV-IL-1 β .

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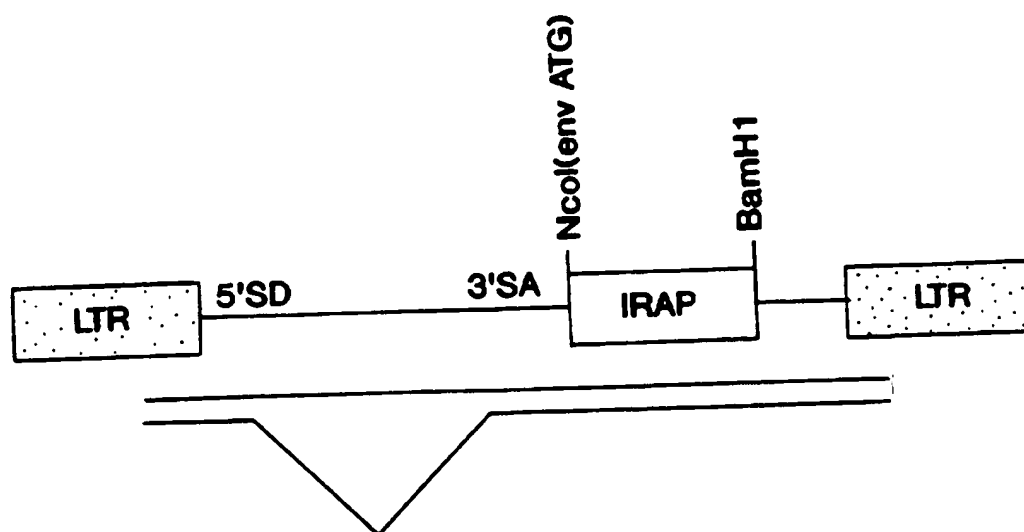


FIG. 1

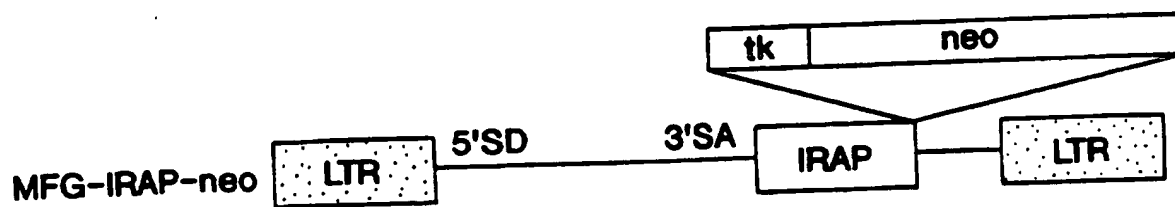


FIG. 2

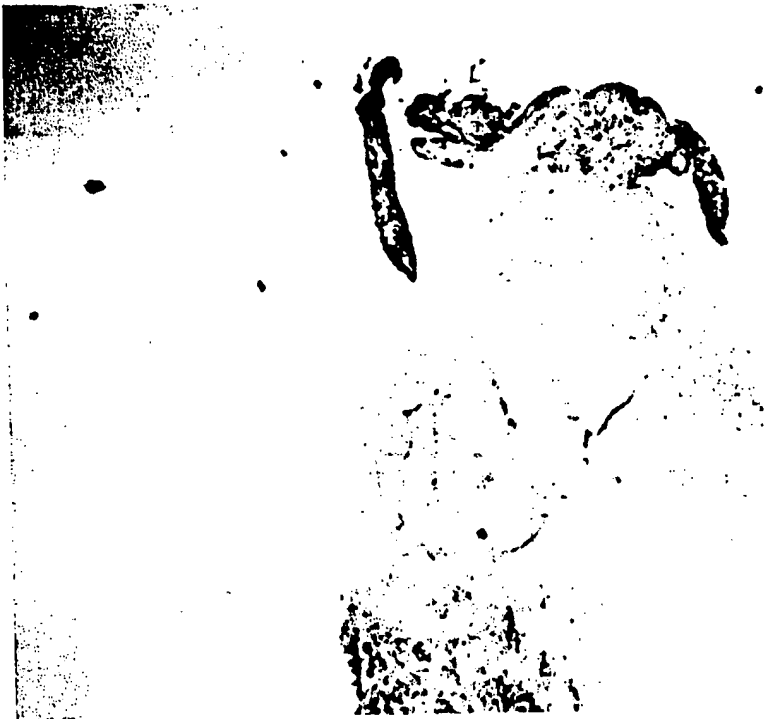


FIG. 3

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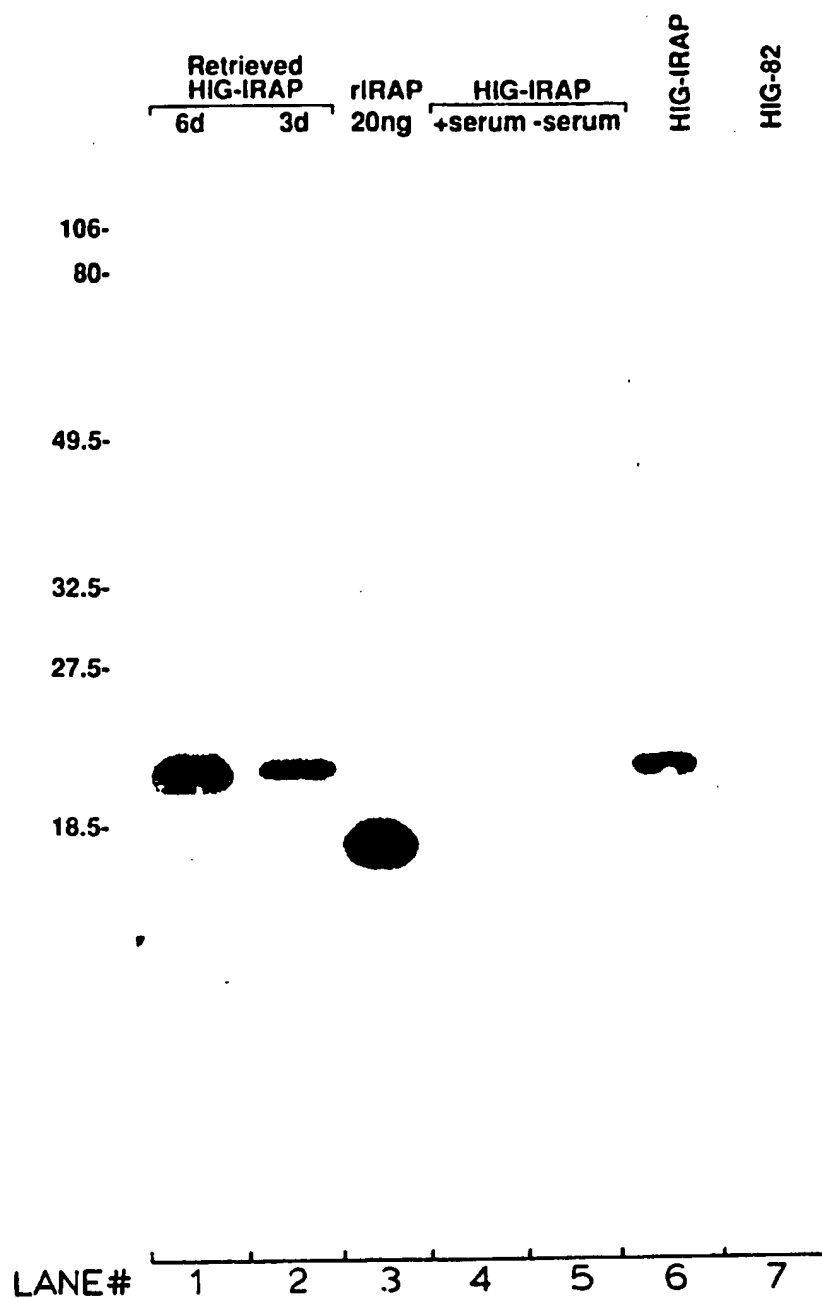


FIG. 4

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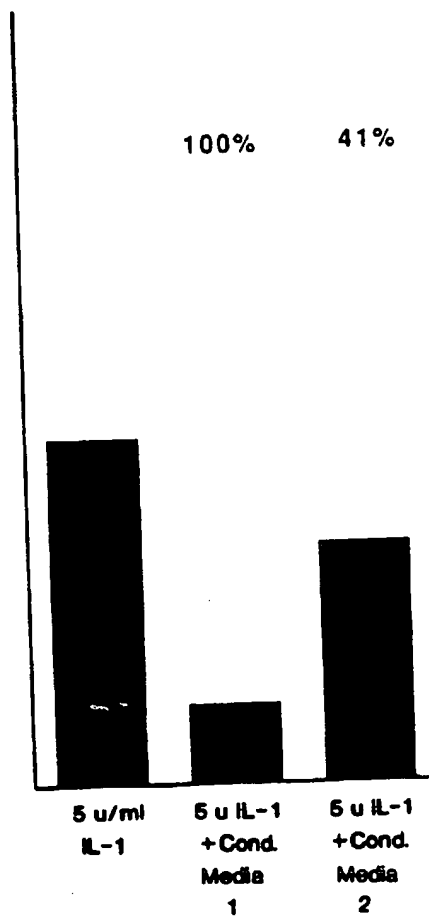
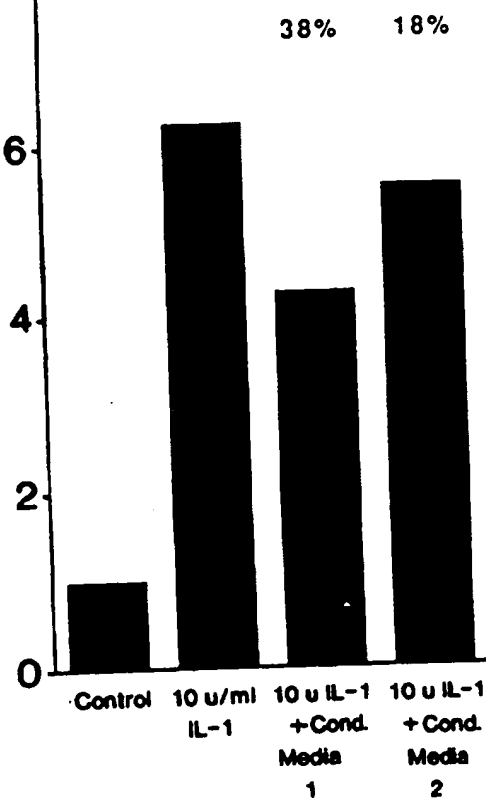
Units Gelatinase/
 10^6 Chondrocytes8
% Inhibition

FIG. 5

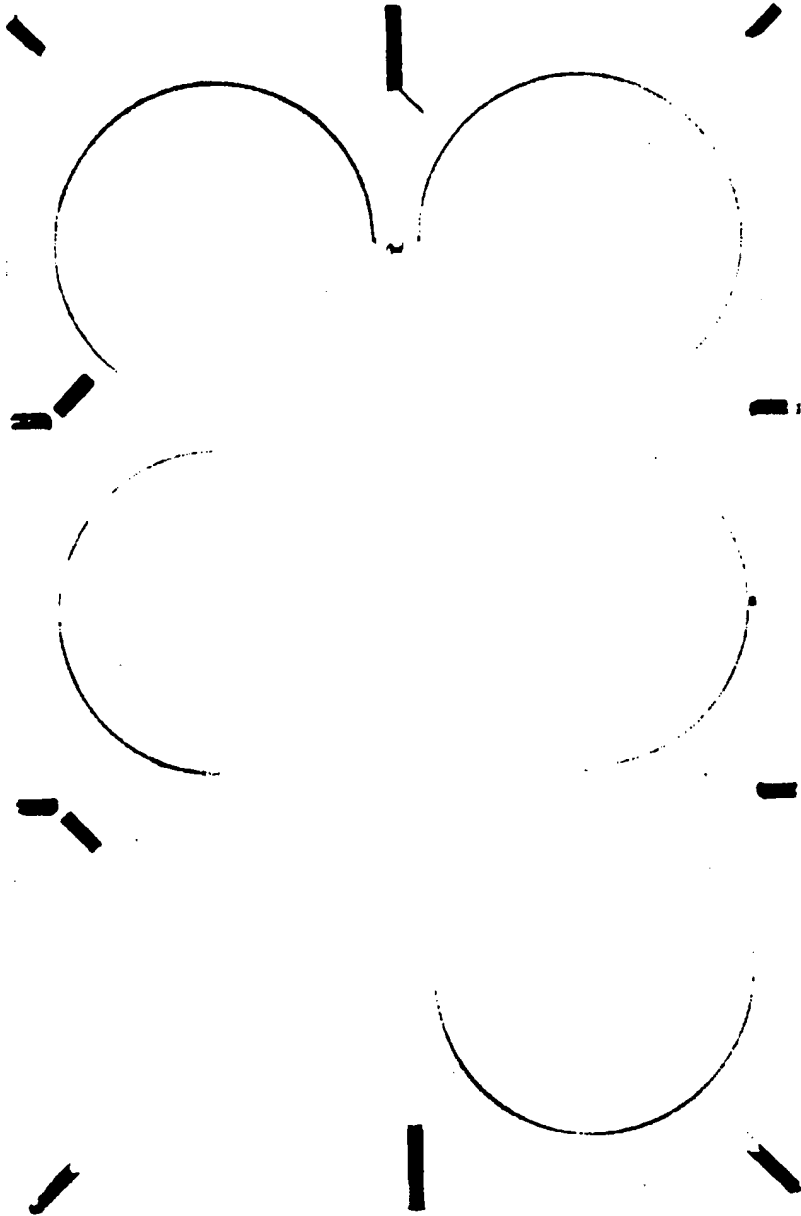
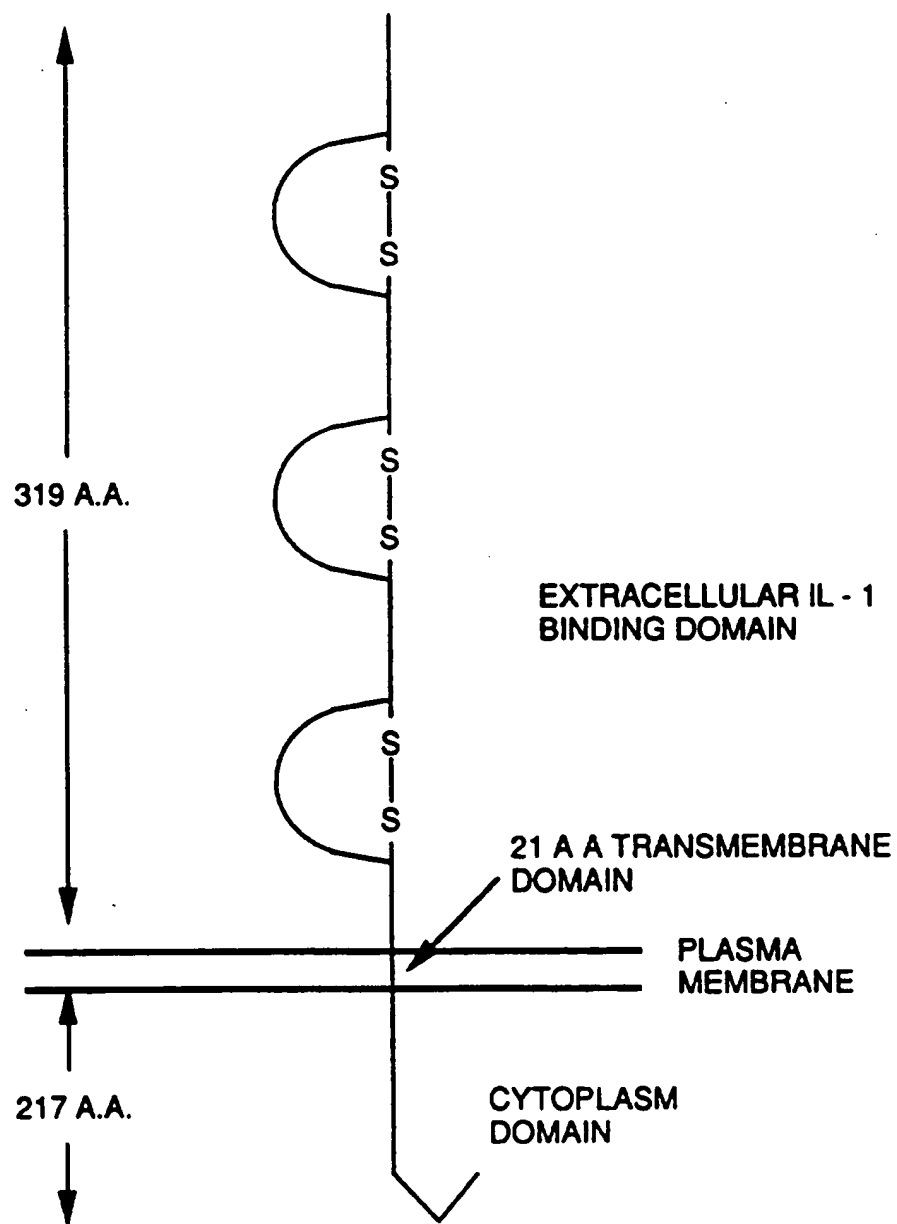


FIG. 6

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**FIG. 7**

[illegible]

FIG. 8A

[illegible]

FIG. 8B

Structure Of The PLJ - ILrec Retroviral Vector
And Partial Restriction Endonuclease Map

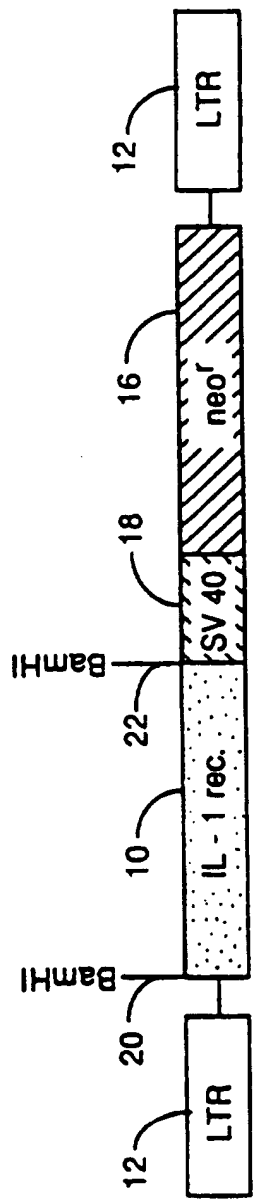


FIG. 9

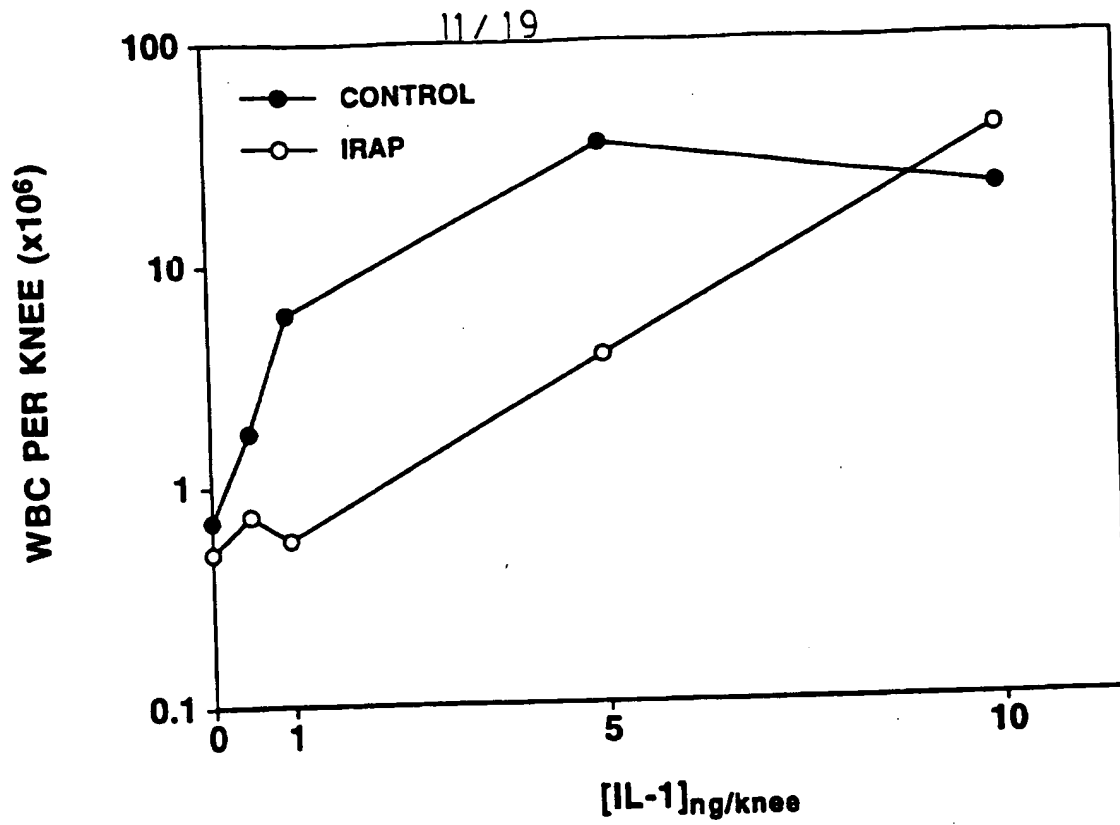
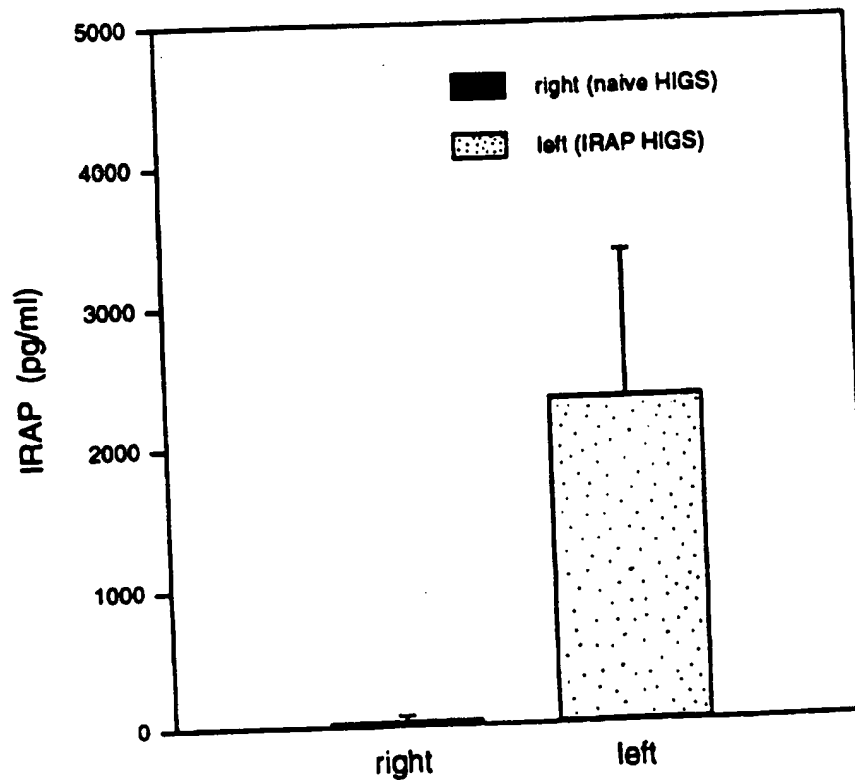


FIG. 10

FIG. 11
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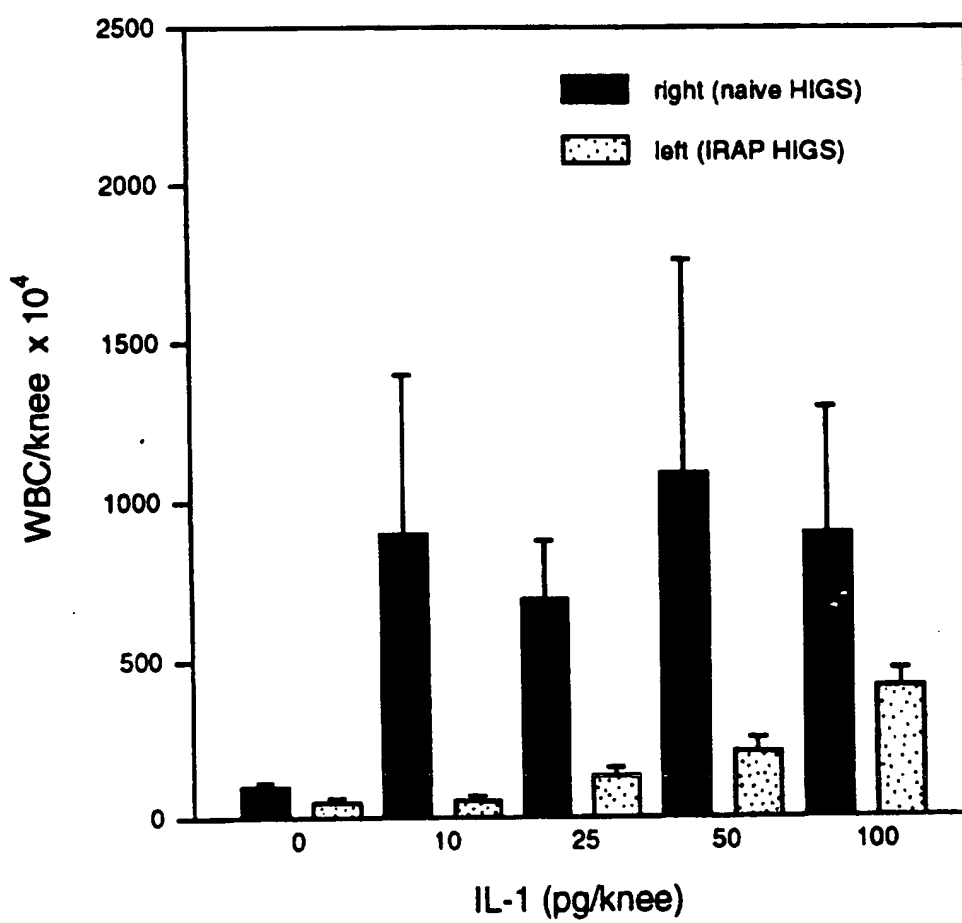


FIG. 12A

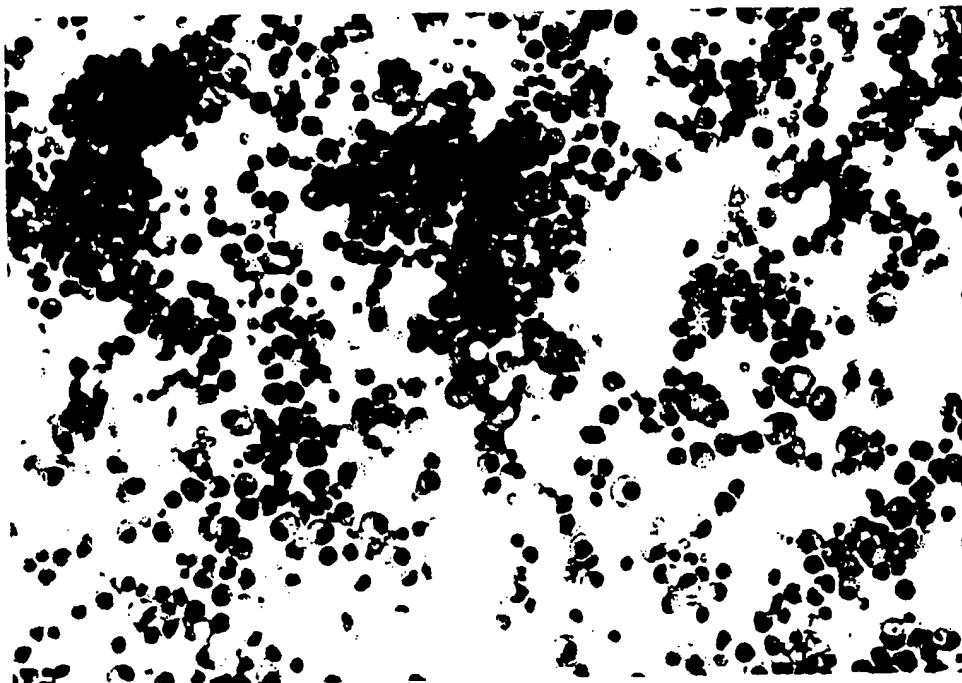


FIG. 12B

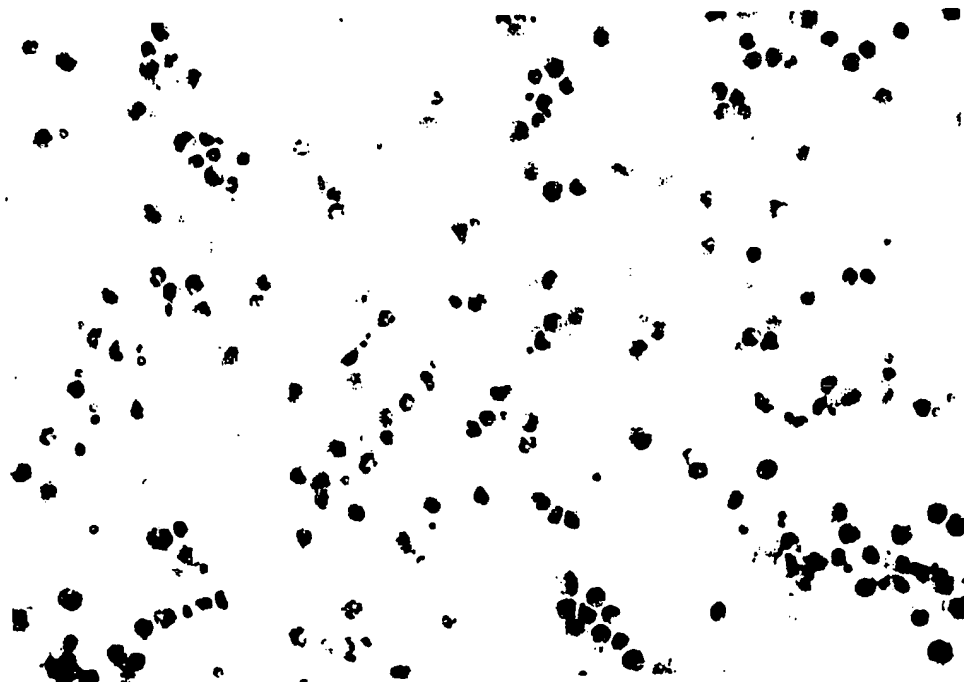


FIG. 12C

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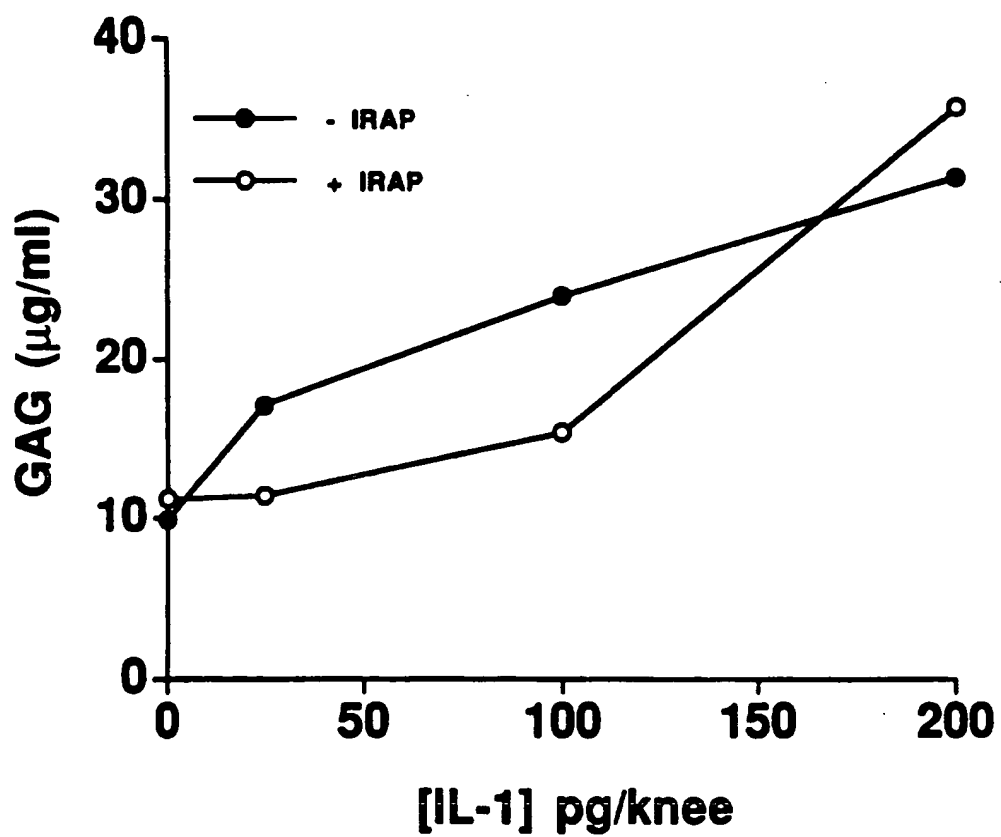


FIG. 13

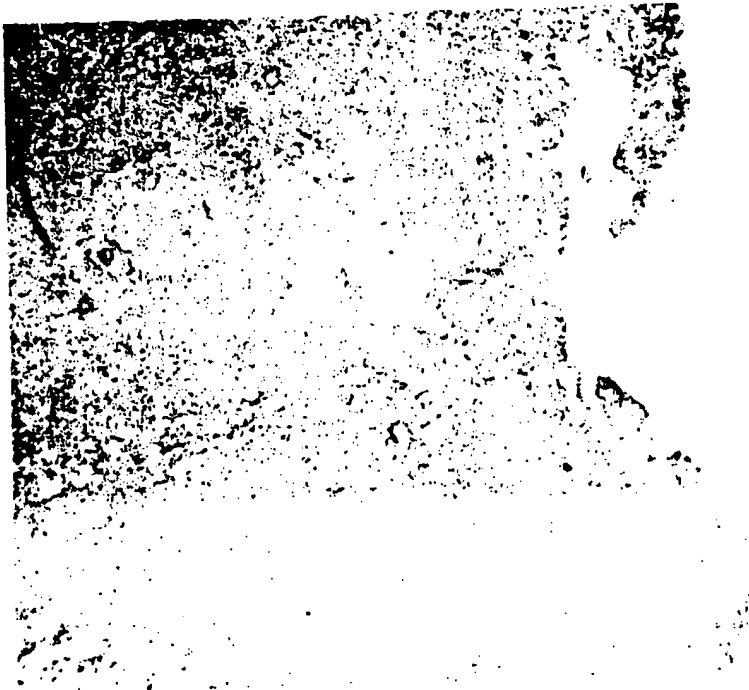


FIG. 14A

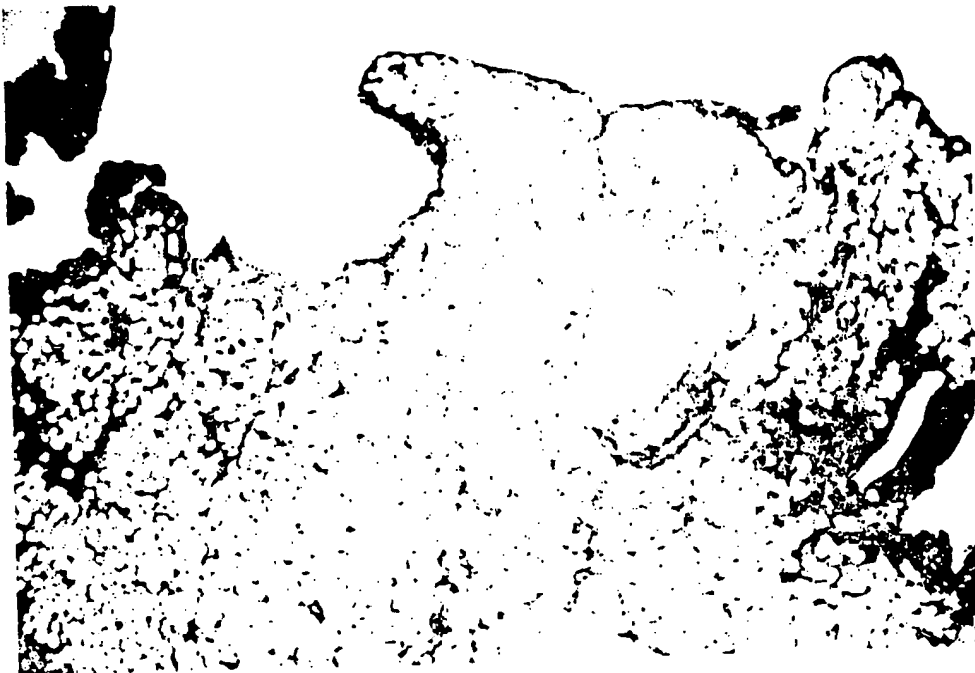


FIG. 14B

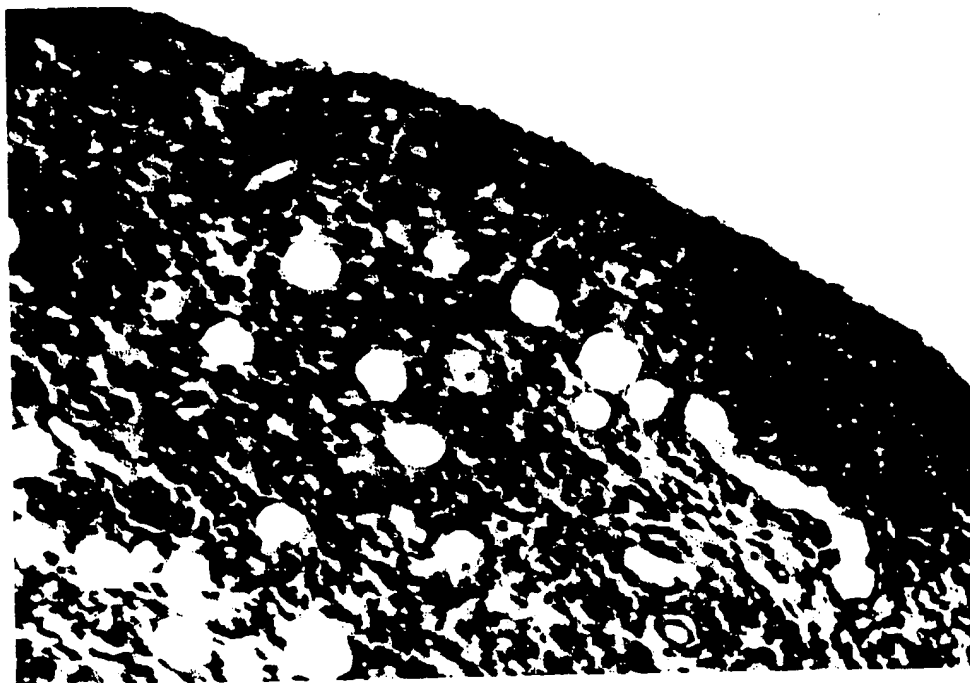


FIG. 14C

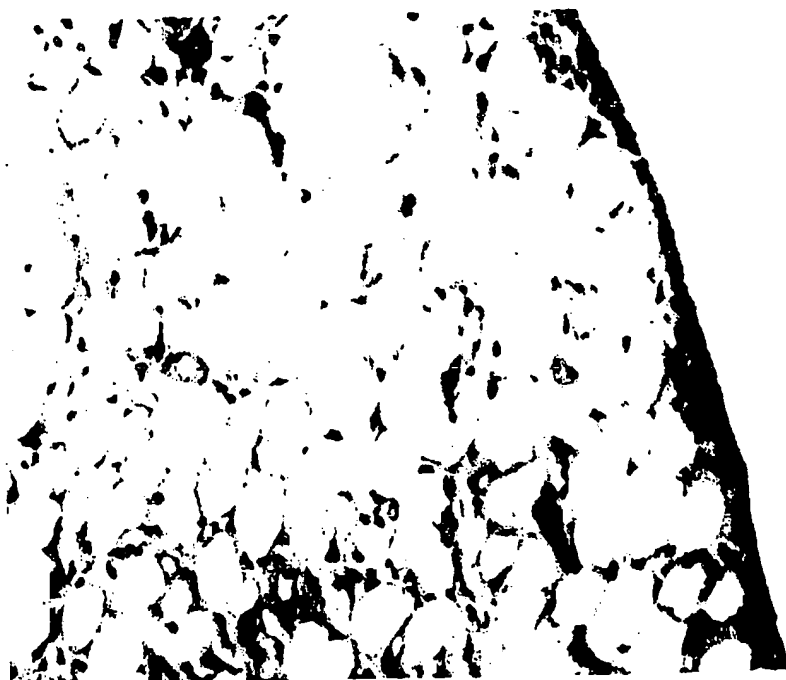


FIG. 14D

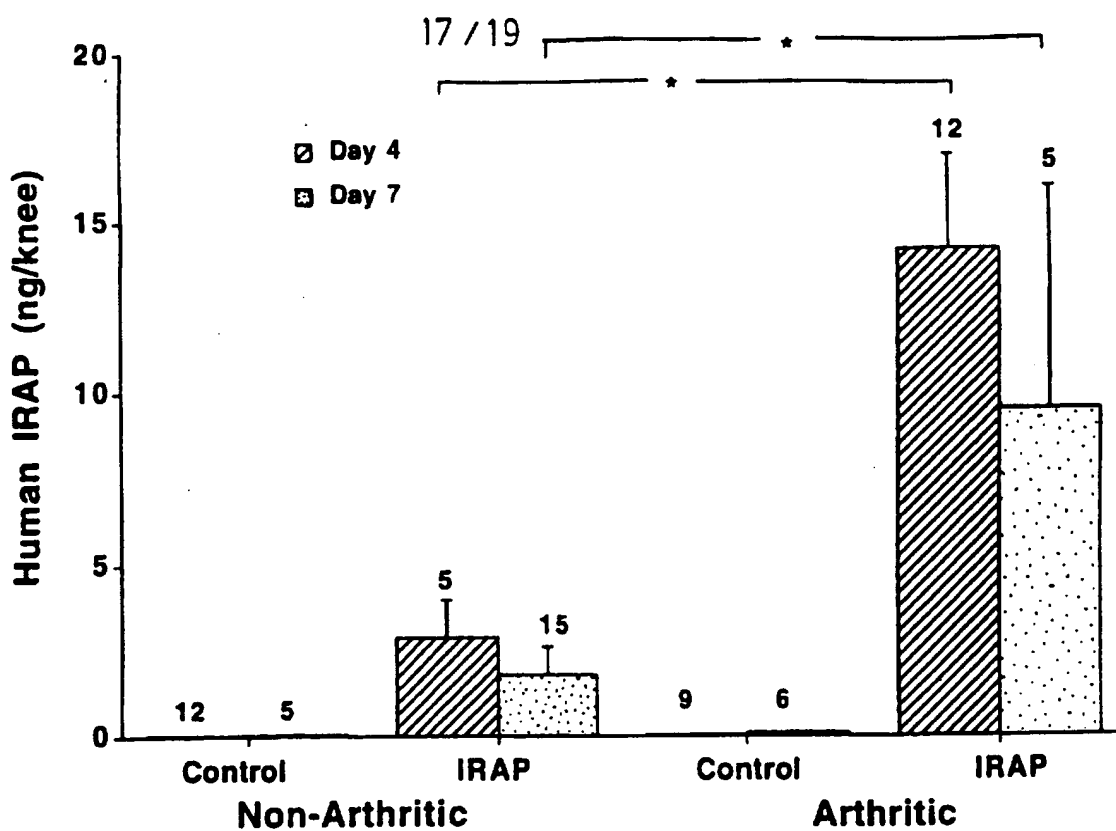


FIG. 15

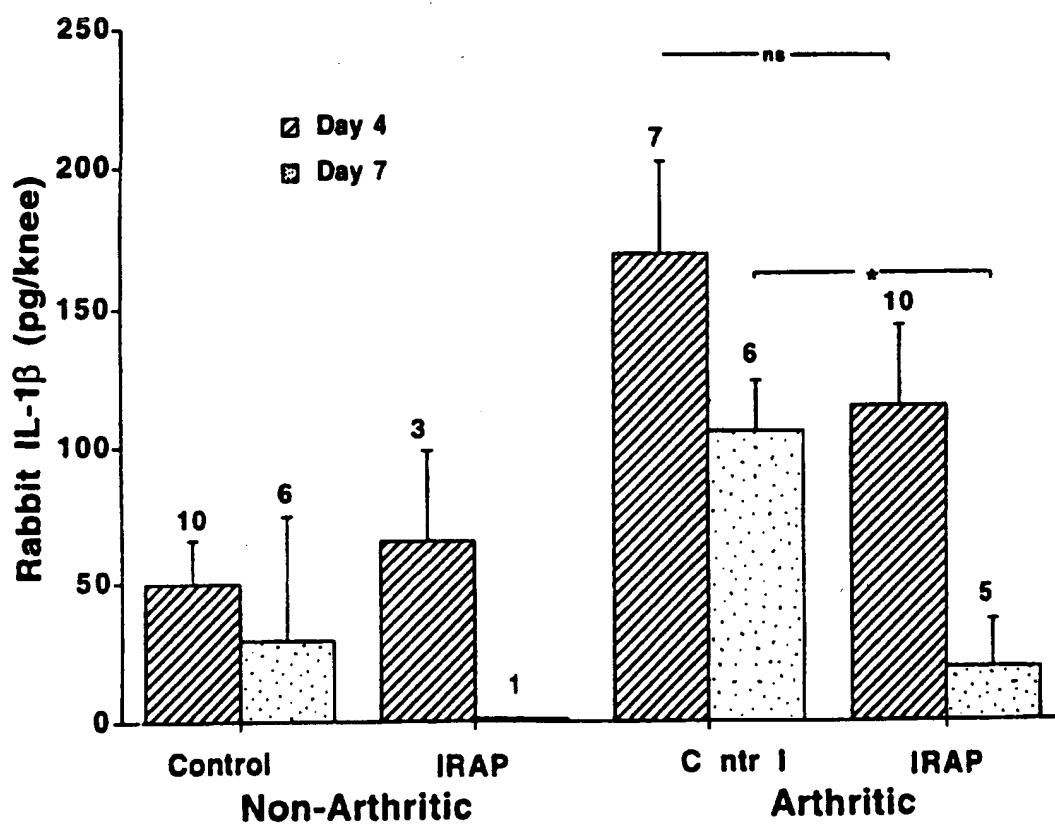


FIG. 16

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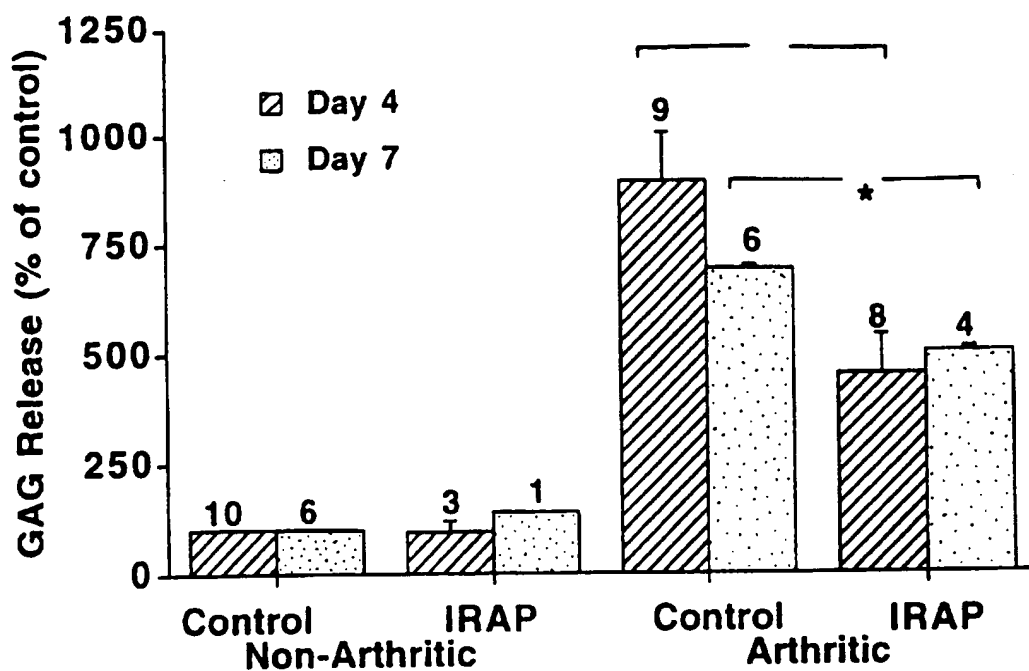


FIG. 17A

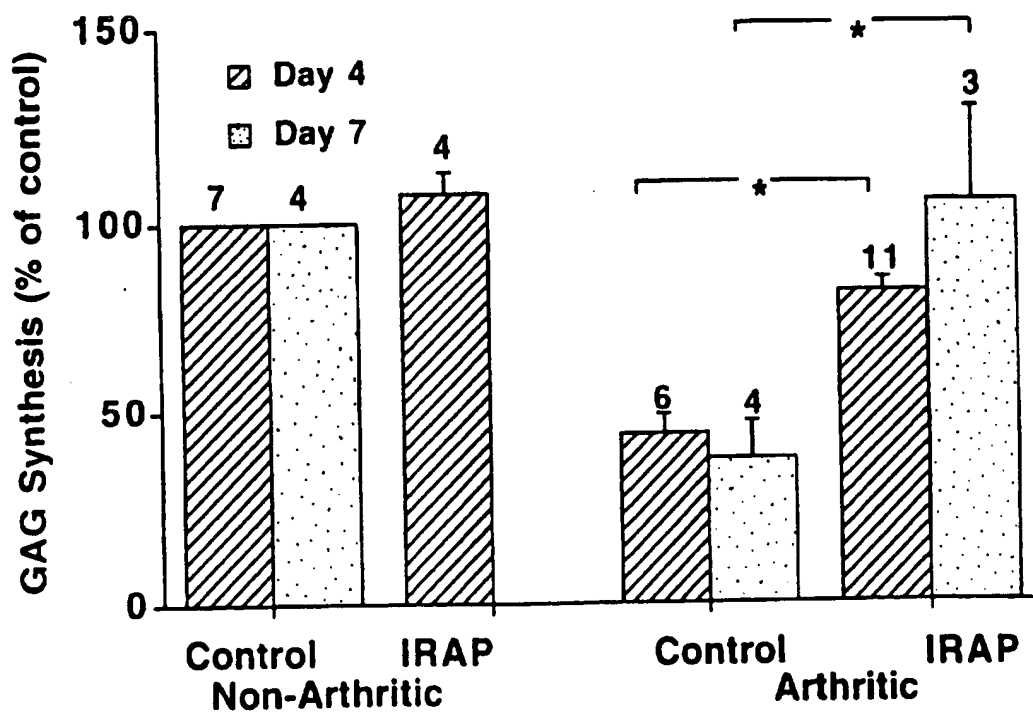


FIG. 17B

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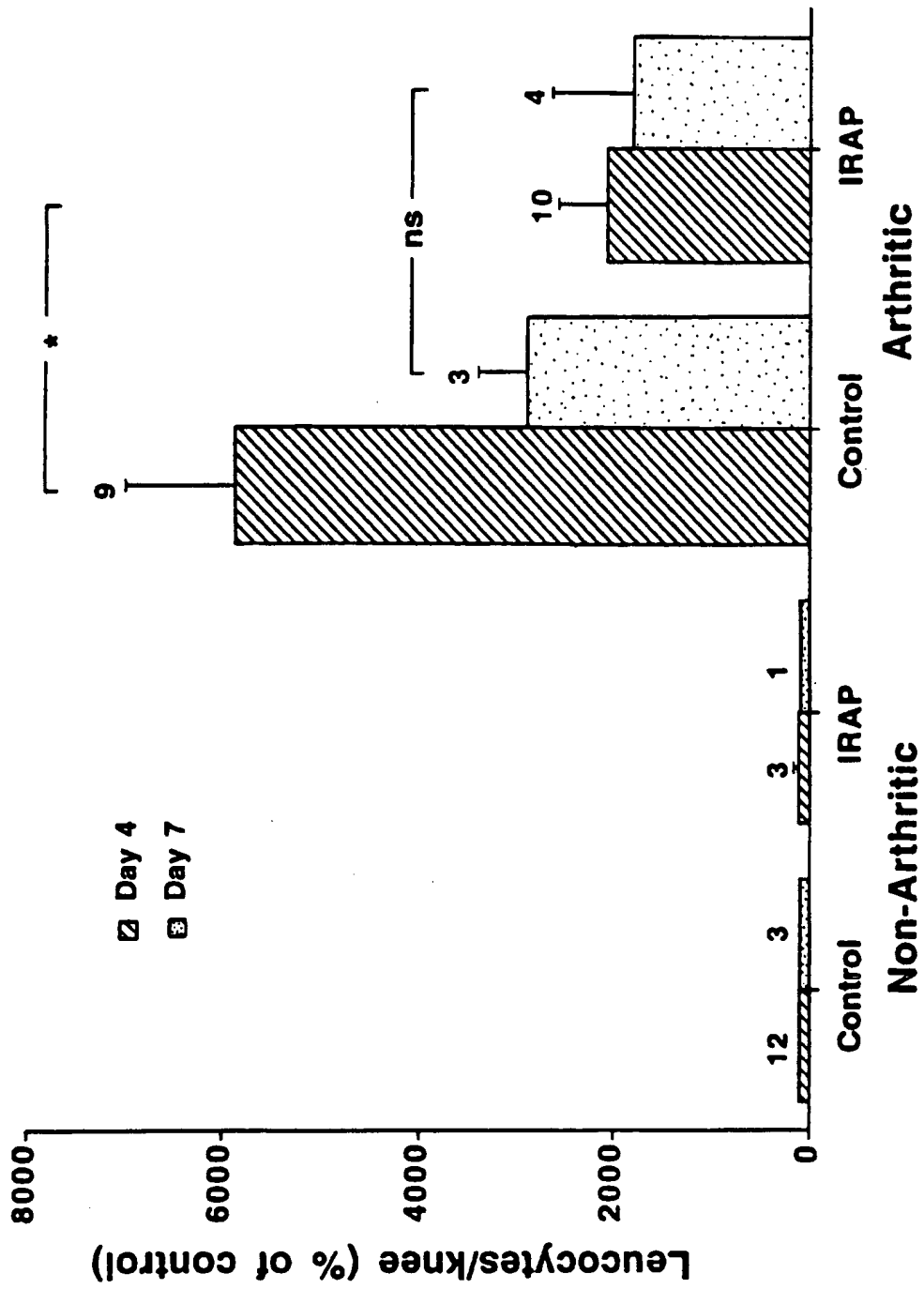


FIG. 18

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US96/01448

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 48/00, 49/00; A01N 63/00

US CL : 514/44; 424/9, 93.1

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/44; 424/9, 93.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, Chemical Abstracts

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Clinical and Experimental Immunology, Volume 96, issued 1994, H. Endo et al, "Experimental Arthritis Induced by Continuous Infusion of IL-8 into Rabbit Knee Joints", pages 31-35, see especially pages 34-35, "Discussion".	27-42

☐

Further documents are listed in the continuation of Box C.

☐

See patent family annex.

* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z*	document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means		
P document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

21 MAY 1996

Date of mailing of the international search report

30 MAY 1996

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Facsimile No. (703) 305-3230

Authorized officer

DEBORAH CROUCH, PH.D.

Telephone No. (703) 308-0196